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(54) **CARBOXYLIC ACID DERIVATIVES, THEIR PREPARATION AND USE**

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See application file for complete search history.

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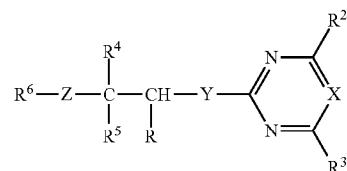
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(57)

**ABSTRACT**

Carboxylic acid derivatives



where R-R<sup>6</sup>, X, Y and Z have the meanings stated in the description, and the preparation thereof, are described. The novel compounds are suitable for controlling diseases.

**23 Claims, No Drawings**

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## CARBOXYLIC ACID DERIVATIVES, THEIR PREPARATION AND USE

Matter enclosed in heavy brackets [ ] appears in the original patent but forms no part of this reissue specification; matter printed in italics indicates the additions made by reissue.

Two (2) reissue applications have been co-filed for the reissue of U.S. Pat. No. 5,932,730. The reissue applications are U.S. Ser. No. 12/481,594 (the present application) and U.S. Ser. No. 12/481,598 (a co-filed reissue application), all of which are co-filed reissues of U.S. Pat. No. 5,932,730.

The present invention relates to novel carboxylic acid derivatives, their preparation and use.

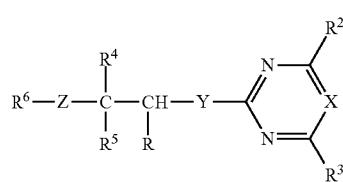
Endothelin is a peptide which is composed of 21 amino acids and is synthesized and released by the vascular endothelium. Endothelin exists in three isoforms, ET-1, ET-2 and ET-3. In the following text, "endothelin" or "ET" signifies one or all isoforms of endothelin. Endothelin is a potent vasoconstrictor and has a potent effect on vessel tone. It is known that this vasoconstriction is caused by binding of endothelin to its receptor (Nature, 332, (1988) 411-415; FEBS Letters, 231, (1988) 440-444 and Biochem. Biophys. Res. Commun., 154, (1988) 868-875).

Increased or abnormal release of endothelin causes persistent vasoconstriction in the peripheral, renal and cerebral blood vessels, which may lead to illnesses. It has been reported in the literature that elevated plasma levels of endothelin were found in patients with hypertension, acute myocardial infarct, pulmonary hypertension, Raynaud's syndrome, atherosclerosis and in the airways of asthmatics (Japan J. Hypertension, 12, (1989) 79, J. Vascular Med. Biology 2, (1990) 207, J. Am. Med. Association 264, (1990) 2868).

Accordingly, substances which specifically inhibit the binding of endothelin to the receptor ought also to antagonize the various abovementioned physiological effects of endothelin and therefore be valuable drugs.

We have found that certain carboxylic acid derivatives are good inhibitors of endothelin receptors.

The invention relates to carboxylic acid derivatives of the formula I



where R is formyl, tetrazole [sic], nitrile [sic], [a COOH group] —COOH or a radical which can be hydrolyzed to —COOH, and the other substituents have the following meanings:

R<sup>2</sup> is hydrogen, hydroxyl, —NH<sub>2</sub>, —NH(C<sub>1</sub>-C<sub>4</sub>-alkyl), —N(C<sub>1</sub>-C<sub>4</sub>-alkyl)<sub>2</sub>, halogen, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy or C<sub>1</sub>-C<sub>4</sub>-alkylthio;

X is nitrogen or CR<sup>14</sup> where R<sup>14</sup> is hydrogen or [C<sub>1-5</sub>] C<sub>1</sub>-C<sub>5</sub>-alkyl, or CR<sup>14</sup> forms together with CR<sup>3</sup> a 5- or 6-membered alkylene or alkenylene ring which can be

substituted by one or two [C<sub>1-4</sub>] C<sub>1</sub>-C<sub>4</sub>-alkyl groups and in which in each case a methylene group can be replaced by oxygen, sulfur, —NH— or —N[C<sub>1-4</sub>](C<sub>1</sub>-C<sub>4</sub>-alkyl)—;

R<sup>3</sup> is hydrogen, hydroxyl, —NH<sub>2</sub>, —NH(C<sub>1</sub>-C<sub>4</sub>-alkyl), —N(C<sub>1</sub>-C<sub>4</sub>-alkyl)<sub>2</sub>, halogen, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy, —NH—O—[C<sub>1-4</sub>] C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-alkylthio or CR<sup>3</sup> is linked to CR<sup>14</sup> as indicated above to give a 5- or 6-membered ring;

R<sup>4</sup> and R<sup>5</sup> (which can be identical or different) are: phenyl or naphthyl, which can be substituted by one or more of the following radicals: halogen, nitro, cyano, hydroxyl, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy, phenoxy, C<sub>1</sub>-C<sub>4</sub>-alkylthio, amino, C<sub>1</sub>-C<sub>4</sub>-alkylamino or C<sub>1</sub>-C<sub>4</sub>-dialkylamino; or phenyl or naphthyl, which are connected together in the ortho positions via a direct linkage, a methylene, ethylene or ethylene group, an oxygen or sulfur atom or an —SO<sub>2</sub>—, —NH— or N-alkyl group, or C<sub>3</sub>-C<sub>7</sub>-cycloalkyl;

R<sup>6</sup> is hydrogen, C<sub>1</sub>-C<sub>8</sub>-alkyl, C<sub>3</sub>-C<sub>6</sub>-alkenyl, C<sub>3</sub>-C<sub>6</sub>-alkynyl or C<sub>3</sub>-C<sub>8</sub>-cycloalkyl, where each of these radicals can be substituted one or more times by: halogen, nitro, cyano, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>3</sub>-C<sub>6</sub>-alkenyloxy, C<sub>3</sub>-C<sub>6</sub>-alkynylloxy, C<sub>1</sub>-C<sub>4</sub>-alkylthio, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy, C<sub>1</sub>-C<sub>4</sub>-alkylcarbonyl, C<sub>1</sub>-C<sub>4</sub>-alkoxycarbonyl, [C<sub>3-8</sub>] C<sub>3</sub>-C<sub>8</sub>-alkylcarbonylalkyl, C<sub>1</sub>-C<sub>4</sub>-alkylamino, di-C<sub>1</sub>-C<sub>4</sub>-alkylamino, phenyl or phenyl or phenoxy which is substituted one or more times, [e.g.] e.g., one to three times, by halogen, [mitro] nitro, cyano, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy or C<sub>1</sub>-C<sub>4</sub>-alkylthio;

phenyl or naphthyl, each of which can be substituted by one or more of the following radicals: halogen, nitro, cyano, hydroxyl, amino, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy, C<sub>1</sub>-C<sub>4</sub>-alkylthio, C<sub>1</sub>-C<sub>4</sub>-alkylamino, C<sub>1</sub>-C<sub>4</sub>-dialkylamino, dioxomethylene [sic] or dioxaethylene [sic]; or

a five- or six-membered heteroaromatic moiety containing one to three nitrogen atoms and/or one sulfur or oxygen atom, which can carry one to four halogen atoms and/or one or two of the following radicals: C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy, C<sub>1</sub>-C<sub>4</sub>-alkylthio, phenyl, phenoxy or phenylcarbonyl, it being possible for the phenyl radicals in turn to carry one to five halogen atoms and/or one to three of the following radicals: C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy and/or C<sub>1</sub>-C<sub>4</sub>-alkylthio;

with the proviso that R<sup>6</sup> can be hydrogen only when Z is not a single bond;

Y is sulfur [or], oxygen or a single bond;

Z is sulfur [or], oxygen or a single bond.

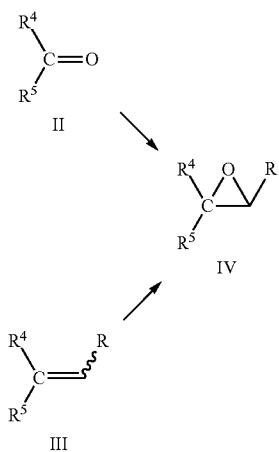
The compounds, and the intermediates for preparing them, such as IV and VI, may have one or more asymmetrical substituted carbon atoms. Such compounds may be in the form of the pure enantiomers or pure diastereomers or a mixture thereof. The use of an enantiomerically pure compound as active substance is preferred.

The invention furthermore relates to the use of the above-mentioned carboxylic acid derivatives for producing drugs, in particular for producing endothelin receptor inhibitors.

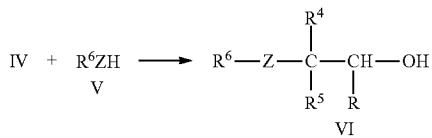
The invention furthermore relates to the preparation of the compounds of the formula IV in enantiomerically pure form. Enantioselective epoxidation of an olefin with two phenyl

substituents is known (J. Org. Chem. 59, 1994, 4378-4380). We have now found, surprisingly, that even ester groups in these systems permit epoxidation in high optical purity.

The preparation of the compounds according to the invention where Z is sulfur or oxygen starts from the epoxides IV, which are obtained in a conventional manner, [e.g.] e.g., as described in J. March, Advanced Organic Chemistry, 2nd ed., 1983, page 862 and page 750, from the ketones II or the olefins III:



Carboxylic acid derivatives of the general formula VI can be prepared by reacting the epoxides of the general formula IV ([e.g.] e.g., with R=ROOR<sup>10</sup> [[sic]]) with alcohols or thiols of the general formula V where R<sup>6</sup> and Z have the meanings stated in claim 1.



To do this, compounds of the general formula IV are heated with compounds of the formula V, in the molar ratio of about 45 1:1 to 1:7, preferably 1 to 3 mole equivalents, to 50-200° C., preferably 80-150° C.

The reaction can also take place in the presence of a diluent. All solvents which are inert toward the reagents used can be used for this purpose.

Examples of such solvents or diluents are water, aliphatic, alicyclic and aromatic hydrocarbons, which may in each case be chlorinated, such as hexane, cyclohexane, petroleum ether, naphtha, benzene, toluene, xylene, methylene chloride, chloroform, carbon tetrachloride, ethyl chloride and trichloroethylene, ethers such as diisopropyl ether, dibutyl ether, methyl tert-butyl ether, propylene oxide, dioxane and tetrahydrofuran, ketones such as acetone, methyl ethyl ketone, methyl isopropyl ketone and methyl isobutyl ketone, nitriles such as acetonitrile and propionitrile, alcohols, such as methanol, ethanol, isopropanol, butanol and ethylene glycol, esters such as ethyl acetate and amyl acetate, amides such as dimethylformamide, dimethylacetamide and N-methylpyrrolidone, sulfoxides and sulfones, such as dimethyl sulfoxide and sulfolane, bases such as pyridine, cyclic ureas such as 1,3-dimethylimidazolidin-2-one and 1,3-dimethyl-3,4,5,6-tetrahydro-2(1H)-pyrimidinone.

The reaction is preferably carried out at a temperature in the range from 0° C. to the boiling point of the solvent or mixture of solvents.

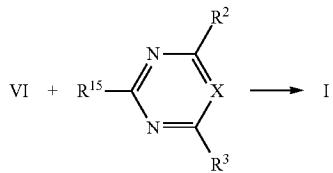
The presence of a catalyst may be advantageous. Suitable catalysts are strong organic and inorganic acids, and Lewis acids. Examples thereof are, inter alia, sulfuric acid, hydrochloric acid, trifluoroacetic acid, p-toluenesulfonic acid, boron trifluoride etherate and titanium(IV) alcoholates.

Compounds of the formula VI where R<sup>4</sup> and R<sup>5</sup> are cycloalkyl can also be prepared by subjecting compounds of the formula VI where R<sup>4</sup> and R<sup>5</sup> are phenyl, naphthyl, or phenyl or naphthyl substituted as described above, to a nuclear hydrogenation.

Compounds of the formula VI can be obtained in enantiomerically pure form by starting from enantiomerically pure compounds of the formula IV and reacting them in the manner described with compounds of the formula V.

It is furthermore possible to obtain enantiomerically pure compounds of the formula VI by carrying out a classical racemate resolution on racemic or diastereomeric compounds of the formula VI using suitable enantiomerically pure bases such as brucine, strychnine, quinine, quinidine, chinchonidine [[sic]], chinchonine [[sic]], yohimbine, morphine, dehydroabietylamine, ephedrine (-), (+), deoxyephedrine (+), (-), threo-2-amino-1-(p-nitrophenyl)-1,3-propanediol (+), (-), threo-2-(N,N-dimethylamino)-1-(p-nitrophenyl)-1,3-propanediol (+), (-), threo-2-amino-1-phenyl-1,3-propanediol (+), (-), α-methylbenzylamine (+), (-), α-(1-naphthyl)ethylamine (+), (-), α-(2-naphthyl)ethylamine (+), (-), aminomethylpinane, N,N-dimethyl-1-phenylethylamine, N-methyl-1-phenylethylamine, 4-nitrophenylethylamine, pseudoephedrine, norephedrine, norpseudoephedrine, amino acid derivatives, peptide derivatives.

The compounds according to the invention where Y is oxygen, and the remaining substituents have the meanings stated under the general formula I, can be prepared, for example, by reacting the carboxylic acid derivatives of the general formula VI where the substituents have the stated meanings with compounds of the general formula VII

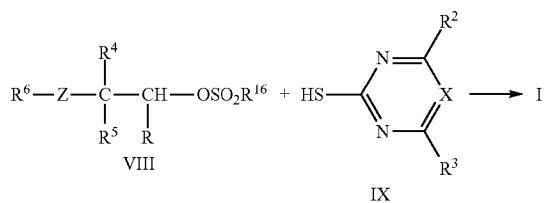


where R<sup>15</sup> is halogen or R<sup>16</sup>-SO<sub>2</sub>-, where R<sup>16</sup> can be C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl or phenyl. The reaction preferably takes place in one of the abovementioned inert diluents with the addition of a suitable base, [i.e.] i.e., of a base which deprotonates the intermediate VI, in a temperature range from room temperature to the boiling point of the solvent.

Compounds of the formula VII are known, some of them can be bought, or they can be prepared in a generally known manner.

It is possible to use as a base an alkali metal or alkaline earth metal hydride such as sodium hydride, potassium hydride or calcium hydride, a carbonate such as an alkali metal carbonate, [e.g.] e.g., sodium or potassium carbonate, an alkali metal or alkaline earth metal hydroxide such as sodium or potassium hydroxide, an organometallic compound such as butyllithium, or an alkali metal amide such as lithium diisopropylamide.

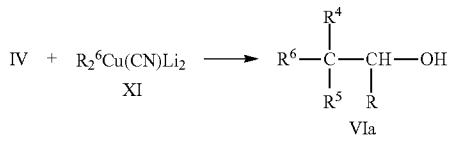
The compounds according to the invention where Y is sulfur, and the remaining substituents have the meanings stated under the general formula I, can be prepared, for example, by reacting carboxylic acid derivatives of the general formula VIII, which can be obtained in a known manner from compounds of the general formula VI and in which the substituents have the abovementioned meanings, with compounds of the general formula IX, where R<sup>2</sup>, R<sup>3</sup> and X have the meanings stated under general formula I.



The reaction preferably takes place in one of the above-mentioned inert diluents with the addition of a suitable base, [i.e.] *i.e.*, a base which deprotonates the intermediate IX, in a temperature range from room temperature to the boiling point of the solvent.

It is possible to use as a base, besides those mentioned above, organic bases such as triethylamine, pyridine, imidazole or diazabicycloundecane [[sic]].

Carboxylic acid derivatives of the formula VIa (z in formula VI=direct linkage) can be prepared by reacting epoxides of the formula IV with cuprates of the formula XI:



The cuprates can be prepared as described in *Tetrahedron Letters* 23, (1982) 3755.

Compounds of the formula I can also be prepared by starting from the corresponding carboxylic acids, [i.e.] *i.e.*, compounds of the formula I where R is COOH, and initially converting these in a conventional manner into an activated form, such as a halide, an anhydride or imidazolide, and then reacting the latter with an appropriate hydroxy compound HOR<sup>10</sup>. This reaction can be carried out in the usual solvents and often requires addition of a base, in which case those mentioned above are suitable. These two steps can also be simplified, for example, by allowing the carboxylic acid to act on the hydroxy compound in the presence of a dehydrating agent such as a carbodiimide.

In addition, it is also possible for compounds of the formula I to be prepared by starting from the salts of the corresponding carboxylic acids, [i.e.] *i.e.*, from compounds of the formula I where R is COR<sup>1</sup> and R<sup>1</sup> is OM, where M can be an alkali metal cation or the equivalent of an alkaline earth metal cation. These salts can be reacted with many compounds of the formula R<sup>1</sup>-A where A is a conventional nucleophilic leaving group, for example halogen such as chlorine, bromine, iodine or aryl- or alkylsulfonyl which is unsubstituted or substituted by halogen, alkyl or haloalkyl, such as toluenesulfonyl and methylsulfonyl, or another equivalent leaving group. Compounds of the formula R<sup>1</sup>-A with a reactive substituent A are known or can be easily obtained with general

expert knowledge. This reaction can be carried out in conventional solvents and advantageously takes place with the addition of a base, in which case those mentioned above are suitable.

5 The radical R in formula I may vary widely. For example, R is a group



where R<sup>1</sup> has the following meanings:

- hydrogen;
- succinylimidoxy [[sic]];
- a five-membered heteroaromatic moiety linked by a nitrogen atom, such as pyrrolyl, pyrazolyl, imidazolyl and triazolyl, which may carry one or two halogen atoms, in particular fluorine and chlorine and/or one or two of the following radicals:

C<sub>1</sub>-C<sub>4</sub>-alkyl such as methyl, ethyl, 1-propyl, 2-propyl, 2-methyl-2-propyl, 2-methyl-1-propyl, 1-butyl, 2-butyl;

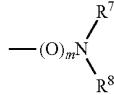
C<sub>1</sub>-C<sub>4</sub>-haloalkyl, in particular C<sub>1</sub>-C<sub>2</sub>-haloalkyl such as fluoromethyl, difluoromethyl, trifluoromethyl, chlorodifluoromethyl, dichlorofluoromethyl, trichloromethyl, 1-fluoroethyl, 2-fluoroethyl, 2,2-difluoroethyl, 2,2,2-trifluoroethyl, 2-chloro-2,2-difluoroethyl, 2,2-dichloro-2-fluoroethyl, 2,2,2-trichloroethyl and pentafluoroethyl;

C<sub>1</sub>-C<sub>4</sub>-haloalkoxy, in particular C<sub>1</sub>-C<sub>2</sub>-haloalkoxy such as difluoromethoxy, trifluoromethoxy, chlorodifluoromethoxy, 1-fluoroethoxy, 2-fluoroethoxy, 2,2-difluoroethoxy, 1,1,2,2-tetrafluoroethoxy, 2,2,2-trifluoroethoxy, 2-chloro-1,1,2-trifluoroethoxy and pentafluoroethoxy, in particular trifluoromethoxy;

C<sub>1</sub>-C<sub>4</sub>-alkoxy such as methoxy, ethoxy, propoxy, 1-methylethoxy, butoxy, 1-methylpropoxy, 2-methylpropoxy, 1,1-dimethylethoxy, in particular methoxy, ethoxy, 1-methylethoxy;

C<sub>1</sub>-C<sub>4</sub>-alkylthio such as methylthio, ethylthio, propylthio, 1-methylethylthio, butylthio, 1-methylpropylthio, 2-methylpropylthio, 1,1-dimethylethylthio, in particular methylthio and ethylthio;

- R<sup>1</sup> is furthermore a radical



where m is 0 or 1 and R<sup>7</sup> and R<sup>8</sup>, which can be identical or different, have the following meanings:

hydrogen;

C<sub>1</sub>-C<sub>8</sub>-alkyl, in particular C<sub>1</sub>-C<sub>4</sub>-alkyl as mentioned above;

C<sub>3</sub>-C<sub>6</sub>-alkenyl such as 2-propenyl, 2-but enyl, 3-but enyl, 1-methyl-2-propenyl, 2-methyl-2-propenyl, 2-pentenyl, 3-pentenyl, 4-pentenyl, 1-methyl-2-but enyl, 2-methyl-2-but enyl, 3-methyl-2-but enyl, 1-methyl-3-but enyl, 2-methyl-3-but enyl, 3-methyl-3-but enyl, 1,1-dimethyl-2-propenyl, 1,2-dimethyl-2-propenyl, 1-ethyl-2-propenyl, 2-hexenyl, 3-hexenyl, 4-hexenyl, 5-hexenyl, 1-methyl-2-pentenyl, 2-methyl-2-pentenyl, 3-methyl-2-pentenyl, 4-methyl-2-pentenyl, 3-methyl-3-pentenyl, 4-methyl-3-pentenyl, 1-me-

thyl-4-pentenyl, 2-methyl-4-pentenyl, 3-methyl-4-pentenyl, 4-methyl-4-pentenyl, 1,1-dimethyl-2-butenyl, 1,1-dimethyl-3-butenyl, 1,2-dimethyl-2-butenyl, 1,2-dimethyl-3-butenyl, 1,3-dimethyl-2-butenyl, 1,3-dimethyl-3-butenyl, 2,2-dimethyl-3-butenyl, 2,3-dimethyl-2-butenyl, 2,3-dimethyl-3-butenyl, 1-ethyl-2-butenyl, 1-ethyl-3-butenyl, 2-ethyl-2-butenyl, 2-ethyl-3-butenyl, 1,1,2-trimethyl-2-propenyl, 1-ethyl-1-methyl-2-propenyl and 1-ethyl-2-methyl-2-propenyl, in particular 2-propenyl, 2-butenyl, 3-methyl-2-butenyl and 3-methyl-2-pentenyl;

$C_3$ - $C_6$ -alkynyl such as 2-propynyl, 2-butyynyl, 3-butyynyl, 1-methyl-2-propynyl, 2-pentynyl, 3-pentynyl, 4-pentynyl, 1-methyl-3-butynyl, 2-methyl-3-butynyl, 1-methyl-2-butynyl, 1,1-dimethyl-2-propynyl, 1-ethyl-2-propynyl, 2-hexynyl, 3-hexynyl, 4-hexynyl, 5-hexynyl, 1-methyl-2-pentynyl, 1-methyl-2-pentynyl, 1-methyl-3-pentynyl, 1-methyl-4-pentynyl, 2-methyl-3-pentynyl, 2-methyl-4-pentynyl, 3-methyl-4-pentynyl, 4-methyl-2-pentynyl, 1,1-dimethyl-2-butynyl, 1,1-dimethyl-3-butynyl, 1,2-dimethyl-3-butynyl, 2,2-dimethyl-3-butynyl, 1-ethyl-2-butynyl, 1-ethyl-3-butynyl, 2-ethyl-3-butynyl and 1-ethyl-1-methyl-2-propynyl, preferably 2-propynyl, 2-butyynyl, 1-methyl-2-propynyl and 1-methyl-2-butynyl, in particular 2-propynyl; or

$C_3$ - $C_8$ -cycloalkyl such as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl [and], cycloheptyl, and cyclooctyl, where these alkyl, cycloalkyl, alkenyl and alkynyl groups can each carry one to five halogen atoms, in particular fluorine or chlorine and/or one or two of the following groups:

$C_1$ - $C_4$ -alkyl,  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -alkylthio,  $C_1$ - $C_4$ -haloalkoxy as mentioned above,  $C_3$ - $C_6$ -alkenyl-<sup>35</sup> loxy,  $C_3$ - $C_6$ -alkenylthio,  $C_3$ - $C_6$ -alkynylloxy,  $C_3$ - $C_6$ -alkynylthio, where the alkenyl and alkynyl constituents present in these radicals preferably have the abovementioned meanings;

$C_1$ - $C_4$ -alkylcarbonyl such as, in particular, methylcarbonyl, ethylcarbonyl, propylcarbonyl, 1-methylethylcarbonyl, butylcarbonyl, 1-methylpropylcarbonyl, 2-methylpropylcarbonyl, 1,1-dimethylethylcarbonyl;

$C_1$ - $C_4$ -alkoxycarbonyl such as methoxycarbonyl, 45 ethoxycarbonyl, propyloxycarbonyl, 1-methylethoxycarbonyl, butyloxycarbonyl, 1-methylpropyloxycarbonyl, 2-methylpropyloxycarbonyl, 1,1-dimethylethoxycarbonyl;

$C_3$ - $C_6$ -alkenylcarbonyl,  $C_3$ - $C_6$ -alkynylcarbonyl,  $C_3$ - $C_6$ -alkenylloxycarbonyl and  $C_3$ - $C_6$ -alkynylloxycarbonyl, where the alkenyl and alkynyl radicals are preferably defined as detailed above;

phenyl, unsubstituted or substituted one or more times, [eg.] e.g., one to three times, by halogen, 55 nitro, cyano,  $C_1$ - $C_4$ -alkyl,  $C_1$ - $C_4$ -haloalkyl,  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -haloalkoxy or  $C_1$ - $C_4$ -alkylthio, such as 2-fluorophenyl, 3-chlorophenyl, 4-bromophenyl, 2-methylphenyl, 3-nitrophenyl, 4-cyanophenyl, 2-trifluoromethylphenyl, 3-methoxyphenyl, 40 4-trifluoroethoxyphenyl, 2-methylthiophenyl, 2,4-dichlorophenyl, 2-methoxy-3-methylphenyl, 2,4-dimethoxyphenyl, 2-nitro-5-cyanophenyl, 2,6-difluorophenyl;

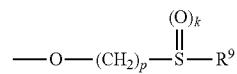
di- $C_1$ - $C_4$ -alkylamino such as, in particular, dimethylamino, dipropylamino, N-propyl-N-methylamino, N-propyl-N-ethylamino, diisopropyl-

lamino, N-isopropyl-N-methylamino, N-isopropyl-N-ethylamino, N-isopropyl-N-propylamino;

$R^7$  and  $[R^8] R^8$  are furthermore phenyl which can be substituted by one or more, [eg.] e.g., one to three, of the following radicals: halogen, nitro, cyano,  $C_1$ - $C_4$ -alkyl,  $C_1$ - $C_4$ -haloalkyl,  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -haloalkoxy or  $C_1$ - $C_4$ -alkylthio, as mentioned above in particular;

or  $R^7$  and  $R^8$  together form a  $C_4$ - $C_7$ -alkylene chain which is closed to form a ring, is unsubstituted or substituted, [eg.] e.g., substituted by  $C_1$ - $C_4$ -alkyl, and may contain a heteroatom selected from the group consisting of oxygen, sulfur or nitrogen, such as  $-(CH_2)_4-$ ,  $-(CH_2)_5-$ ,  $-(CH_2)_6-$ ,  $-(CH_2)_7-$ ,  $-(CH_2)_2-O-(CH_2)_2-$ ,  $-CH_2-S-(CH_2)_3-$ ,  $-(CH_2)_2-O-(CH_2)_3-$ ,  $-NH-(CH_2)_3-$ ,  $-CH_2-NH-(CH_2)_2-$ ,  $-CH_2-CH=CH-CH_2-$ ,  $-CH=CH-(CH_2)_3-$ ;

e)  $R^1$  is furthermore a group



where  $k$  is 0, 1 and 2; [ ];

$p$  is 1, 2, 3 and 4; and

$R^9$  is  $C_1$ - $C_4$ -alkyl,  $C_1$ - $C_4$ -haloalkyl,  $C_3$ - $C_6$ -alkenyl,  $C_3$ - $C_6$ -alkynyl or unsubstituted or substituted phenyl, as mentioned above in particular [ ].

f)  $R^1$  is furthermore a radical OR<sup>10</sup>, where R<sup>10</sup> is:

hydrogen, the cation of an alkali metal such as lithium, sodium, potassium or the cation of an alkaline earth metal such as calcium, magnesium and barium or an environmentally compatible organic ammonium ion such as tertiary  $C_1$ - $C_4$ -alkylammonium or the ammonium ion;

$C_3$ - $C_8$ -cycloalkyl as mentioned above, which may carry one to three  $C_1$ - $C_4$ -alkyl groups;

$C_1$ - $C_8$ -alkyl such as, in particular, methyl, ethyl, propyl, 1-methylethyl, butyl, 1-methylpropyl, 2-methylpropyl, 1,1-dimethylethyl, pentyl, 1-methylbutyl, 2-methylbutyl, 3-methylbutyl, 1,2-dimethylpropyl, 1,1-dimethylpropyl, 2,2-dimethylpropyl, 1-ethylpropyl, hexyl, 1-methylpentyl, 2-methylpentyl, 3-methylpentyl, 4-methylpentyl, 1,2-dimethylbutyl, 1,3-dimethylbutyl, 2,3-dimethylbutyl, 1,1-dimethylbutyl, 2,2-dimethylbutyl, 3,3-dimethylbutyl, 1,1,2-trimethylpropyl, 1,2,2-trimethylpropyl, 1-ethylbutyl, 2-ethylbutyl, 1-ethyl-2-methylpropyl, which can carry one to five halogen atoms, in particular fluorine and chlorine and/or one of the following radicals:

$C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -alkylthio, cyano,  $C_1$ - $C_4$ -alkylcarbonyl,  $C_3$ - $C_8$ -cycloalkyl,  $C_1$ - $C_4$ -alkoxycarbonyl, phenyl, phenoxy or phenylcarbonyl, where the aromatic radicals in turn can carry in each case one to five halogen atoms and/or one to three of the following radicals: nitro, cyano,  $C_1$ - $C_4$ -alkyl,  $C_1$ - $C_4$ -haloalkyl,  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -haloalkoxy and/or  $C_1$ - $C_4$ -alkylthio, as mentioned above in particular;

[a]  $C_1$ - $C_8$ -alkyl as mentioned above, which can carry one to five halogen atoms, in particular fluorine and/or chlorine, and carries one of the following

radicals: a 5-membered heteroaromatic moiety containing one to three nitrogen atoms, or a 5-membered heteroaromatic moiety containing a nitrogen atom and an oxygen or sulfur atom, which can carry one to four halogen atoms and/or one or two of the following radicals:

nitro, cyano,  $C_1$ - $C_4$ -alkyl,  $C_1$ - $C_4$ -haloalkyl,  $C_1$ - $C_4$ -alkoxy, phenyl,  $C_1$ - $C_4$ -haloalkoxy and/or  $C_1$ - $C_4$ -alkylthio. Particular mention may be made of: 1-pyrazolyl, 3-methyl-1-pyrazolyl, 4-methyl-1-pyrazolyl, 3,5-dimethyl-1-pyrazolyl, 3-phenyl-1-pyrazolyl, 4-phenyl-1-pyrazolyl, 4-chloro-1-pyrazolyl, 4-bromo-1-pyrazolyl, 1-imidazolyl, 1-benzimidazolyl, 1,2,4-triazol-1-yl, 3-methyl-1,2,4-triazol-1-yl, 5-methyl-1,2,4-triazol-1-yl, 1-benzotriazolyl, 3-isopropyl-5-isoxazolyl, 3-methyl-5-isoxazolyl, 2-oxazolyl, 2-thiazolyl, 2-imidazolyl, 3-ethyl-5-isoxazolyl, 3-phenyl-5-isoxazolyl, 3-tert-butyl-5-isoxazolyl;

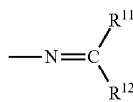
[a]  $C_2$ - $C_6$ -alkyl [group] which carries one of the following radicals in position 2:  $C_1$ - $C_4$ -alkoxyimino,  $C_3$ - $C_6$ -alkynyoxyimino,  $C_3$ - $C_6$ -haloalkenyloxyimino or benzyloxyimino; or

[a]  $C_3$ - $C_6$ -alkenyl or  $C_3$ - $C_6$ -alkynyl [group], it being possible for these groups in turn to carry one to five halogen atoms;

$R^{10}$  is furthermore a phenyl radical which can carry one to five halogen atoms and/or one to three of the following radicals: nitro, cyano,  $C_1$ - $C_4$ -alkyl,  $C_1$ - $C_4$ -haloalkyl,  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -haloalkoxy and/or  $C_1$ - $C_4$ -alkylthio, as mentioned above in particular;

a 5-membered heteroaromatic moiety which is linked via a nitrogen atom, contains one to three nitrogen atoms and can carry one or two halogen atoms and/or one or two of the following radicals:  $C_1$ - $C_4$ -alkyl,  $C_1$ - $C_4$ -haloalkyl,  $C_1$ - $C_4$ -alkoxy, phenyl,  $C_1$ - $C_4$ -haloalkoxy and/or  $C_1$ - $C_4$ -alkylthio. Particular mention may be made of: 1-pyrazolyl, 3-methyl-1-pyrazolyl, 4-methyl-1-pyrazolyl, 3,5-dimethyl-1-pyrazolyl, 3-phenyl-1-pyrazolyl, 4-phenyl-1-pyrazolyl, 4-chloro-1-pyrazolyl, 4-bromo-1-pyrazolyl, 1-imidazolyl, 1-benzimidazolyl, 1,2,4-triazol-1-yl, 3-methyl-1,2,4-triazol-1-yl, 5-methyl-1,2,4-triazol-1-yl, 1-benzotriazolyl, 3,4-dichloro-1-imidazolyl;

$R^{10}$  is furthermore a group



where  $R^{11}$  and  $R^{12}$ , which can be identical or different, are:

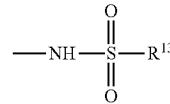
$C_1$ - $C_8$ -alkyl,  $C_3$ - $C_6$ -alkenyl,  $C_3$ - $C_6$ -alkynyl,  $C_3$ - $C_8$ -cycloalkyl, it being possible for these radicals to carry a  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -alkylthio and/or an unsubstituted or substituted phenyl radical, as mentioned above in particular; phenyl which can be substituted by one or more,

[eg.] e.g., one to three, of the following radicals: halogen, nitro, cyano,  $C_1$ - $C_4$ -alkyl,  $C_1$ - $C_4$ -haloalkyl,  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -haloalkoxy or

$C_1$ - $C_4$ -alkylthio, where these radicals are, in particular, those mentioned above;

or  $R^{11}$  and  $R^{12}$  together form a  $C_3$ - $C_{12}$ -alkylene chain which can carry one to three  $C_1$ - $C_4$ -alkyl groups and contain a heteroatom from the group consisting of oxygen, sulfur and nitrogen, as mentioned in particular for  $R^7$  and  $R^8$  [.]

g)  $R^1$  is furthermore a radical

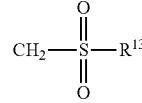


where  $R^{13}$  is:

$C_1$ - $C_4$ -alkyl,  $C_3$ - $C_6$ -alkenyl,  $C_3$ - $C_6$ -alkynyl,  $C_3$ - $C_8$ -cycloalkyl as mentioned above in particular, it being possible for these radicals to carry a  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -alkylthio and/or a phenyl radical as mentioned above; or

phenyl, unsubstituted or substituted, in particular as mentioned above [.]

h)  $R^1$  is a radical



where  $R^{13}$  has the abovementioned meaning.

R can furthermore be:

tetrazole [[sic]] or nitrile [[sic]].

In respect of the biological effect, preferred carboxylic acid derivatives of the general formula I, both as pure enantiomers and pure diastereomers or as mixture thereof, are those where the substituents have the following meanings:

$R^2$  is hydrogen, hydroxyl,  $N(C_1$ - $C_4$ -alkyl)<sub>2</sub>, [the]  $C_1$ - $C_4$ -alkyl,  $C_1$ - $C_4$ -haloalkyl,  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -alkylthio groups and halogen atoms mentioned in detail for  $R^1$ , especially chlorine, methyl, methoxy, ethoxy, difluoromethoxy, trifluoromethoxy;

$X$  is nitrogen or  $CR^{14}$  where

$R^{14}$  is hydrogen or alkyl, or  $CR^{14}$  forms together with  $CR^3$  a 4- to 5-membered alkylene or alkenylene ring in which, in each case, a methylene group can be replaced by oxygen or sulfur, such as  $—CH_2—CH_2—O—$ ,  $—CH=CH—O—$ ,  $—CH_2—CH_2—CH_2—O—$ ,  $—CH=CH—CH_2O—$ , in particular hydrogen,  $—CH_2—CH_2—O—$ ,  $—CH(CH_3)—CH(CH_3)—O—$ ,  $—C(CH_3)=C(CH_3)—O—$ ,  $—CH=C(CH_3)—O—$  or  $—C(CH_3)=C(CH_3)—S—$ ;

$R^3$  [the] is hydrogen, hydroxyl,  $N(C_1$ - $C_4$ -alkyl)<sub>2</sub>,  $C_1$ - $C_4$ -alkyl,  $C_1$ - $C_4$ -haloalkyl,  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -haloalkoxy,  $C_1$ - $C_4$ -alkylthio groups and halogen atoms mentioned for  $R^1$ , especially chlorine, methyl, methoxy, ethoxy, difluoromethoxy, trifluoromethoxy or  $R^3$  is linked to  $R^{14}$  as mentioned above to give a 5- or 6-membered ring;

$R^4$  and  $R^5$  are phenyl or naphthyl, which can be substituted by one or more, [eg.] e.g., one to three, of the following radicals: halogen, nitro, cyano, hydroxyl, mercapto, amino,  $C_1$ - $C_4$ -alkyl,  $C_1$ - $C_4$ -haloalkyl,  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -haloalkoxy,  $C_1$ - $C_4$ -alkylthio,  $C_1$ - $C_4$ -alkylamino, di- $C_1$ - $C_4$ -alkylamino,  $C_1$ - $C_4$ -alkylcarbonyl,  $C_1$ - $C_4$ -

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alkoxycarbonyl; phenyl or naphthyl, which are connected together in the ortho positions by a direct linkage, a methylene, ethylene or ethenylene group, an oxygen or sulfur atom or an  $-\text{SO}_2-$ ,  $-\text{NH}-$  [or] N-alkyl group, or  $\text{C}_3\text{-C}_7\text{-cycloalkyl}$ ;

$\text{R}^6$  is  $\text{C}_1\text{-C}_8\text{-alkyl}$ ,  $\text{C}_3\text{-C}_6\text{-alkenyl}$ ,  $\text{C}_3\text{-C}_6\text{-alkynyl}$  or  $\text{C}_3\text{-C}_8\text{-cycloalkyl}$  as mentioned above in particular, it being possible for these radicals in each case to be substituted one or more times by: halogen, hydroxyl, nitro, cyano,  $\text{C}_1\text{-C}_4\text{-alkoxy}$ ,  $\text{C}_3\text{-C}_6\text{-alkenylloxy}$ ,  $\text{C}_3\text{-C}_6\text{-alkynylloxy}$ ,  $\text{C}_1\text{-C}_4\text{-alkylthio}$ ,  $\text{C}_1\text{-C}_4\text{-haloalkoxy}$ ,  $\text{C}_1\text{-C}_4\text{-alkylcarbonyl}$ , hydroxycarbonyl,  $\text{C}_1\text{-C}_4\text{-alkoxycarbonyl}$ ,  $\text{C}_1\text{-C}_4\text{-alkylamino}$ , di- $\text{C}_1\text{-C}_4\text{-alkylamino}$  or unsubstituted or substituted phenyl or phenoxy, as mentioned above in particular;

phenyl or naphthyl, which can be substituted by one or more of the following radicals: halogen, nitro, cyano, hydroxyl, amino,  $\text{C}_1\text{-C}_4\text{-alkyl}$ ,  $\text{C}_1\text{-C}_4\text{-haloalkyl}$ ,  $\text{C}_1\text{-C}_4\text{-alkoxy}$ ,  $\text{C}_1\text{-C}_4\text{-haloalkoxy}$ , phenoxy,  $\text{C}_1\text{-C}_4\text{-alkylthio}$ ,  $\text{C}_1\text{-C}_4\text{-[akylamino [sic]] alkylamino}$  or  $\text{C}_1\text{-C}_4\text{-dialkylamino}$ , as mentioned in particular for  $\text{R}^7$  and  $\text{R}^4$ ; or

a five- or six-membered heteroaromatic moiety which contains one to three nitrogen atoms and/or one sulfur or oxygen atom and which can carry one to four halogen atoms and/or one or two of the following radicals:  $\text{C}_1\text{-C}_4\text{-alkyl}$ ,  $\text{C}_1\text{-C}_4\text{-haloalkyl}$ ,  $\text{C}_1\text{-C}_4\text{-alkoxy}$ ,  $\text{C}_1\text{-C}_4\text{-haloalkoxy}$ ,  $\text{C}_1\text{-C}_4\text{-alkylthio}$ , phenyl, phenoxy or phenylcarbonyl, it being possible for the phenyl radicals in turn to carry one to five halogen atoms and/or one to three of the following radicals:  $\text{C}_1\text{-C}_4\text{-alkyl}$ ,  $\text{C}_1\text{-C}_4\text{-haloalkyl}$ ,  $\text{C}_1\text{-C}_4\text{-alkoxy}$ ,  $\text{C}_1\text{-C}_4\text{-haloalkoxy}$  and/or  $\text{C}_1\text{-C}_4\text{-alkylthio}$ , as mentioned for  $\text{R}^4$  in particular;

$\text{Y}$  is sulfur, oxygen or a single bond;

$\text{Z}$  is sulfur, oxygen,  $-\text{SO}-$ ,  $-\text{SO}_2-$  or a single bond.

Particularly preferred compounds of the formula I, both as pure enantiomers and pure diastereomers or as mixture thereof, are those in which the substituents have the following meanings:

$\text{R}^2$  is  $\text{C}_1\text{-C}_4\text{-alkyl}$ , or  $\text{C}_1\text{-C}_4\text{-alkoxy}$ ;

$\text{X}$  is nitrogen or  $\text{CR}^{14}$ , where

$\text{R}^{14}$  is hydrogen or alkyl, or  $\text{CR}^{14}$  forms together with  $\text{CR}^3$  a 4- or 5-membered alkylene or alkenylene ring such as  $-\text{CH}_2\text{-CH}_2\text{-CH}_2-$ ,  $-\text{CH}=\text{CH}-\text{CH}_2-$ , in which in each case a methylene group can be replaced by oxygen or sulfur, such as  $-\text{CH}_2\text{-CH}_2\text{-O}-$ ,  $-\text{CH}=\text{CH}-\text{O}-$ ,  $-\text{CH}_2\text{-CH}_2\text{-CH}_2\text{-O}-$ ,  $-\text{CH}=\text{CH}-\text{CH}_2\text{O}-$ , in particular hydrogen,  $-\text{CH}_2\text{-CH}_2\text{-O}-$ ,  $-\text{CH}(\text{CH}_3)\text{-CH}(\text{CH}_3)\text{-O}-$ ,  $-\text{C}(\text{CH}_3)\text{-C}(\text{CH}_3)\text{-O}-$ ,  $-\text{CH}=\text{C}(\text{CH}_3)\text{-O}-$  or  $-\text{C}(\text{CH}_3)\text{-C}(\text{CH}_3)\text{-S}-$ ;

$\text{R}^3$  [the] is  $\text{C}_1\text{-C}_4\text{-alkyl}$ ,  $\text{C}_1\text{-C}_4\text{-alkoxy}$ ,  $\text{C}_1\text{-C}_4\text{-alkylthio}$  groups mentioned for  $\text{R}^1$ , or  $\text{R}^3$  is linked to  $\text{R}^{14}$  as mentioned above to give a 5- or 6-membered ring;

$\text{R}^4$  and  $\text{R}^5$  are phenyl (identical or different) which can be substituted by one or more, [eg.] e.g., one to three, of the following radicals: halogen, nitro, hydroxyl,  $\text{C}_1\text{-C}_4\text{-alkyl}$ ,  $\text{C}_1\text{-C}_4\text{-alkoxy}$ ,  $\text{C}_1\text{-C}_4\text{-alkylthio}$ ; or

$\text{R}^4$  and  $\text{R}^5$  are phenyl groups which are connected together in the ortho positions by a direct linkage, a methylene, ethylene or ethenylene group, an oxygen or sulfur atom or an  $\text{SO}_2$ ,  $\text{NH}$  or N-alkyl group; or

$\text{R}^4$  and  $\text{R}^5$  are  $\text{C}_3\text{-C}_7\text{-cycloalkyl}$ ;

$\text{R}^6$  is  $\text{C}_1\text{-C}_8\text{-alkyl}$ ,  $\text{C}_3\text{-C}_6\text{-alkenyl}$  or  $\text{C}_3\text{-C}_8\text{-cycloalkyl}$ , it being possible for these radicals in each case to be sub-

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stituted one or more times by: halogen, hydroxyl, nitro, cyano,  $\text{C}_1\text{-C}_4\text{-alkoxy}$ ,  $\text{C}_3\text{-C}_6\text{-alkenylloxy}$ ,  $\text{C}_1\text{-C}_4\text{-alkylthio}$ ; or

$\text{R}^6$  is phenyl or naphthyl, which can be substituted by one or more of the following radicals: halogen, nitro, cyano, hydroxyl, amino,  $\text{C}_1\text{-C}_4\text{-alkyl}$ ,  $\text{C}_1\text{-C}_4\text{-haloalkyl}$ ,  $\text{C}_1\text{-C}_4\text{-alkoxy}$ ,  $\text{C}_1\text{-C}_4\text{-alkylthio}$ ,  $\text{C}_1\text{-C}_4\text{-[akylamino [sic]] or C}_1\text{-C}_4\text{-dialkylamino}$ ; or

$\text{R}^6$  is a five- or six-membered heteroaromatic moiety which contains a nitrogen atom and/or a sulfur or oxygen atom and which can carry one to four halogen atoms and/or one or two of the following radicals:  $\text{C}_1\text{-C}_4\text{-alkyl}$ ,  $\text{C}_1\text{-C}_4\text{-haloalkyl}$ ,  $\text{C}_1\text{-C}_4\text{-alkoxy}$ ,  $\text{C}_1\text{-C}_4\text{-alkylthio}$ , phenyl, phenoxy or phenylcarbonyl, it being possible for the phenyl radicals in turn to carry one to five halogen atoms and/or one to three of the following radicals:  $\text{C}_1\text{-C}_4\text{-alkyl}$ ,  $\text{C}_1\text{-C}_4\text{-haloalkyl}$ ,  $\text{C}_1\text{-C}_4\text{-alkoxy}$  and/or  $\text{C}_1\text{-C}_4\text{-alkylthio}$ ;

$\text{Y}$  is sulfur, oxygen or a single bond;

$\text{Z}$  is sulfur, oxygen,  $-\text{SO}-$ ,  $-\text{SO}_2-$  or a single bond.

The compounds of the present invention provide a novel therapeutic potential for the treatment of hypertension, pulmonary hypertension, myocardial infarct, angina pectoris, acute kidney failure, renal insufficiency, cerebral vasospasms, cerebral ischemia, subarachnoid hemorrhages, migraine, asthma, atherosclerosis, endotoxic shock, endotoxin-induced organ failure, intravascular coagulation, restenosis after angioplasty, benign prostate hyperplasia, or hypertension or kidney failure caused by ischemia or intoxication.

The good effect of the compounds can be shown in the following tests:

Receptor binding studies

35 Cloned human  $\text{ET}_A$  receptor-expressing CHO cells and guinea pig cerebellar membranes with  $>60\%$   $\text{ET}_B$  compared with  $\text{ET}_A$  receptors were used for binding studies.

The  $\text{ET}_A$  receptor-expressing CHO cells were grown in  $\text{F}_{12}$  medium containing 10% fetal calf serum, 1% glutamine, 100 U/ml penicillin and 0.2% streptomycin (Gibco BRL, Gaithersburg, Md., USA).

After 48 h, the cells were washed with PBS and incubated with 0.05% trypsin-containing PBS for 5 min. Neutralization was then carried out with  $\text{F}_{12}$  medium, and the cells were collected by centrifugation at 300xg. To [lyse] lye the cells, the pellet was briefly washed with lysis buffer (5 mM Tris-HCl, pH 7.4 with 10% glycerol) and then incubated at a concentration of 107 cells/ml of lysis buffer at 4° C. for 30 min. The membranes were centrifuged at 20,000xg for 10 min, and the pellet was stored in liquid nitrogen.

Guinea pig cerebella were homogenized in a Potter-Elvehjem homogenizer and [lacuna] obtained by differential centrifugation at 1000xg for 10 min and repeated centrifugation of the supernatant at 20,000xg for 10 min.

Binding assays

For the  $\text{ET}_A$  and  $\text{ET}_B$  receptor binding assay, the membranes were suspended in incubation buffer (50 mM Tris-HCl, pH 7.4 with 5 mM MnCl<sub>2</sub>, 40  $\mu\text{g}/\text{ml}$  bacitracin and 0.2% BSA) at a concentration of 50  $\mu\text{g}$  of protein per assay mixture and incubated with 25 pM [<sup>125</sup>I]<sup>-</sup> $\text{ET}_1$  ( $\text{ET}_A$  receptor assay) or 25 pM [<sup>125</sup>I]<sup>-</sup> $\text{RZ}_3$  ( $\text{ET}_B$  receptor assay) in the presence and absence of test substance at 25° C. The nonspecific binding was determined using [10<sup>-7</sup>] 10<sup>-7</sup> M  $\text{ET}_1$ . After 30 min, the free and bound radioligand were separated by filtration through GF/B glass fiber filters (Whatman, England) on a Skatron cell collector (Skatron, Lier, Norway) and the filters were washed with ice-cold Tris-HCl buffer, pH

7.4 with 0.2% BSA. The radioactivity collected on the filters was quantified using a Packard 2200 CA liquid scintillation counter.

Functional in vitro assay system to look for endothelin receptor (subtype A) antagonists

This assay system is a functional, cell-based assay for endothelin receptors. When certain cells are stimulated with endothelin 1 (ET1) they show an increase in the intracellular calcium concentration. This increase can be measured in intact cells loaded with calcium-sensitive dyes.

1-Fibroblasts which had been isolated from rats and in which an endogenous endothelin receptor of the A subtype had been detected were loaded with the fluorescent dye **[Fura 2-an] Fura 2-am** as follows: after trypsinization, the cells were resuspended in buffer A (120 mM NaCl, 5 mM KCl, 1.5 mM MgCl<sub>2</sub>, 1 mM CaCl<sub>2</sub>, 25 mM HEPES, 10 mM glucose, pH 7.4) to a density of  $2 \times 10^6$ /ml and incubated with Fura 2-am (2  $\mu$ M), Pluronics F-127 (0.04%) **[und]** and DMSO (0.2%) at 37°C. in the dark for 30 min. The cells were then washed twice with buffer A and resuspended at  $2 \times 10^6$ /ml.

The fluorescence signal from  $2 \times 10^5$  cells per ml with Ex/Em 380/510 was recorded continuously at 30°C. The test substances and, after an incubation time of 3 min, ET1 **[lacauna]** to the cells, the maximum change in the fluorescence was determined. The response of the cells to ET1 without previous addition of a test substance was used as control and was set equal to 100%.

Testing of ET antagonists in vivo

Male SD rats weighting 250-300 g were anesthetized with amobarbital, **[artificially]** *artificially* ventilated, vagotomized and pithed. The carotid artery and jugular vein were **[cathetized]** **[sic]** *catheterized*.

In control animals, intravenous administration of 1  $\mu$ g/kg ET1 led to a distinct rise in blood pressure which persisted for a lengthy period.

The test animals received an i.v. injection of the test compounds (1 ml/kg) 5 min before the administration of ET1. To determine the ET-antagonistic properties, the rise in blood pressure in the test animals was compared with that in the control animals.

Endothelin-1-induced sudden death in mice

The principle of the test is the inhibition of the sudden heart death caused in mice by endothelin, which is probably induced by constriction of the coronary vessels, by pretreatment with endothelin receptor antagonists. Intravenous injection of 10 nmol/kg endothelin in a volume of 5 ml/kg of body weight results in death of the animals within a few minutes.

The lethal endothelin-1 dose is checked in each case on a small group of animals. If the test substance is administered intravenously, the endothelin-1 injection which was lethal in the reference group usually takes place 5 min thereafter. With other modes of administration, the times before administration are extended, where appropriate up to several hours.

The survival rate is recorded, and effective doses which protect 50% of the animals (ED 50) from endothelin-induced heart death for 24 h or longer are determined.

Functional test on vessels for endothelin receptor antagonists

Segments of rabbit aorta are, after an initial tension of 2 g and a relaxation time of 1 h in Krebs-Henseleit solution at 37°C. and pH 7.3-7.4, first induced to contract with K<sup>+</sup>. After washing out, an endothelin dose-effect plot up to the maximum is constructed.

Potential endothelin antagonists are administered to other preparations of the same vessel 15 min before starting the endothelin dose-effect plot. The effects of the endothelin are

calibrated as a % of the K<sup>+</sup>-induced contraction. Effective endothelin antagonists result in a shift to the right in the endothelin dose-effect plot.

The compounds according to the invention can be administered orally or parenterally (subcutaneously, intravenously, intramuscularly, **[intraperitoneally]** *intraperitoneally*) in a conventional way. Administration can also take place with vapors or sprays through the nasopharyngeal space.

The dosage depends on the age, condition and weight of the patient and on the mode of administration. The daily dose of active substance is, as a rule, about 0.5-50 mg/kg of body weight on oral administration and about 0.1-10 mg/kg of body weight on parenteral administration.

The novel compounds can be used in conventional solid or liquid pharmaceutical forms, **[e.g.]** *e.g.*, as uncoated or (film-) coated tablets, capsules, powders, granules, suppositories, solutions, ointments, creams or sprays. These are produced in a conventional way. The active substances can for this purpose be processed with conventional pharmaceutical aids such as tablet binders, fillers, preservatives, tablet disintegrants, flow regulators, plasticizers, wetting agents, dispersants, emulsifiers, solvents, release-slowng agents, antioxidants and/or propellant gases (cf. H. Sucker et al.: *Pharmazeutische Technologie*, Thieme-Verlag, Stuttgart, 1991). The administration forms obtained in this way normally contain from 0.1 to 90% by weight of the active substance.

### Synthesis examples

#### Example 1

Methyl 2-hydroxy-3-methoxy-3,3-diphenylpropionate  
5 g (19.6 mmol) of methyl 3,3-diphenyl-2,3-epoxypropionate were dissolved in 50 ml of absolute methanol and, at 0°C., 0.1 ml of boron trifluoride etherate was added. The mixture was stirred at 0°C. for 2 h and at room temperature for a further 12 h. The solvent was distilled out, the residue was taken up in ethyl acetate, washed with sodium bicarbonate solution and water and dried over magnesium sulfate. After removal of the solvent by distillation there remained 5.5 g (88%) of a pale yellow oil.

#### Example 2

Methyl 2-hydroxy-3-phenoxy-3,3-diphenylpropionate  
5 g (19.6 mmol) of methyl 3,3-diphenyl-2,3-epoxypropionate and 5.6 g (60 mmol) of phenol were heated together at 100°C. for 6 h. Removal of the excess phenol by distillation under high vacuum and purification of the residue by chromatography on silica gel with hexane/ethyl acetate mixtures resulted in 4.9 g (77%) of a pale yellow oil.

#### Example 3

Methyl 2-(4,6-dimethoxy-pyrimidin-2-yloxy)-3-methoxy-3,3-diphenylpropionate  
2.86 g (10 mmol) of methyl 2-hydroxy-3-methoxy-3,3-diphenylpropionate were dissolved in 40 ml of dimethylformamide, and 0.3 g (12 mmol) of sodium hydride was added. The mixture was stirred for 1 h and then 2.2 g (10 mmol) of 4,6-dimethoxy-2-methylsulfonylpyrimidine were added. After stirring at room temperature for 24 h, cautious hydrolysis was carried out with 10 ml of water, the pH was adjusted to 5 with acetic acid, and the solvent was removed by distillation under high vacuum. The residue was taken up in 100 ml of ethyl acetate, washed with water and dried over magne-

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sium sulfate, and the solvent was distilled out. The residue was mixed with 10 ml of ether, and the resulting precipitate was filtered off with suction. After drying, 3.48 g (82%) of a white powder remained.

Melting point 81° C.

## Example 4

## 2-(4,6-Dimethoxy-pyrimidin-2-yloxy)-3-methoxy-3,3-diphenylpropionic acid

2.12 g (5 mmol) of methyl 2-(4,6-dimethoxy-pyrimidin-2-yloxy)-3-methoxy-3,3-diphenylpropionate were dissolved in 50 ml of dioxane, 10 ml of 1N KOH solution were added, and the mixture was stirred at 100° C. for 3 h. The solution was diluted with 300 ml of water and extracted with ethyl acetate to remove unreacted ester. The aqueous phase was then adjusted to pH 1-2 with dilute hydrochloric acid and extracted with ethyl acetate. After drying over magnesium sulfate and removal of the solvent by distillation, the residue was mixed with an ether/hexane mixture, and the precipitate which formed was filtered off with suction. After drying, 1.85 g (90%) of a white powder remained.

Melting point 167° C.

## Example 5

## 2-(4,6-Dimethoxy-2-pyrimidinyloxy)-3-methoxy-3,3-diphenyl sodium [sic] propionate

1.68 g (4 mmol) of 2-(4,6-dimethoxy-2-pyrimidinyloxy)-3-methoxy-3,3-diphenylpropionic acid are dissolved in 4 ml of 1N NaOH+100 ml of water. The solution is freeze-dried, and the sodium salt of the carboxylic acid used is obtained quantitatively.

10 g (34.9 mmol) of methyl 2-hydroxy-3-methoxy-3,3-diphenylpropionate were dissolved in 50 ml each of methanol and glacial acetic acid, 1 ml of RuO(OH)<sub>2</sub> in dioxane was added, and hydrogenation was carried out with H<sub>2</sub> in an autoclave at 100° C. under 100 bar for 30 h. The catalyst was filtered off, the mixture was concentrated, mixed with ether and washed with NaCl solution, and the organic phase was dried and concentrated. 10.1 g of methyl 3,3-dicyclohexyl-2-hydroxy-3-methoxypropionate were obtained as an oil.

## Example 7

## Methyl 2-[(4,6-dimethoxy-pyrimidin-2-yl)thio]-3-methoxy-3,3-diphenylpropionate [sic]

7.16 g (25 mmol) of methyl 2-hydroxy-3-methoxy-3,3-diphenylpropionate were dissolved in 50 ml of dichloromethane, 3 g (30 mmol) of triethylamine were added, and 3.2 g (28 mmol) of methanesulfonyl chloride were added dropwise while stirring. The mixture was stirred at room temperature for 2 h, washed with water, dried over magnesium sulfate and concentrated under reduced pressure. The residue was taken up in DMF and added dropwise at 0° C. to a suspension of 12.9 g (75 mmol) of 4,6-dimethoxypyrimidine-2-thiol and 8.4 g (100 mmol) of sodium bicarbonate in 100 ml of DMF. After stirring at room temperature for 2 h and at 60° C. for a further 2 h, the mixture was poured into 1 liter of ice-water, and the resulting precipitate was filtered off with suction. After drying, 3.19 g (29%) of a white powder remained.

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## Example 8

## Methyl 2-hydroxy-3,3-diphenylbutyrate

1.5 g (5.9 mmol) of methyl 3,3-diphenyl-2,3-epoxypropionate dissolved in 10 ml of absolute ether were added dropwise to a cup-rate solution which had been prepared from 635 mg (7 mmol) of copper(I) cyanide dissolved in 10 ml of absolute ether and 8.14 ml (13 mmol) of a 1.6 normal methyl lithium solution and had been cooled to -78° C. The solution was stirred at -78° C. for 1 h and then allowed to warm to room temperature. It was subsequently diluted with 100 ml of ether and 100 ml of water, and the ether phase was washed with dilute citric acid and with sodium bicarbonate solution and dried over magnesium sulfate. The crude product was purified by chromatography on silica gel with cyclohexane/ethyl acetate mixtures to result in 250 mg (16%) of a pale yellow oil.

## Example 9

## 2-Hydroxy-3-methoxy-3,3-diphenylpropionic acid

91.11 g (0.5 mol) of benzophenone and 45.92 g (0.85 mol) of sodium methoxide were suspended in 150 ml of methyl tert-butyl ether (MTB) at room temperature. After cooling to -100° C., 92.24 g (0.85 mol) of methyl chloroacetate were added in such a way that the internal temperature rose to 40° C. while continuing to cool in a bath at -10° C. The mixture was then stirred without cooling at the autogenous temperature for one hour. After addition of 250 ml of water and brief stirring, the aqueous phase was separated off. The MTB phase was washed with 250 ml of dilute sodium chloride solution. After the solvent had been changed to methanol (250 ml), a solution of 1 g of p-toluenesulfonic acid in 10 ml of methanol was added at room temperature. The mixture was stirred at autogenous temperature for one hour and then heated to reflux. While distilling out the methanol, 400 g of a 10% strength sodium hydroxide solution was added dropwise, and finally 60 ml of water were added. The methanol was distilled out until the bottom temperature reached 97° C. After cooling to 55° C., 190 ml of MTB were added and the mixture was acidified to pH 2 with about 77 ml of concentrated HCl. After cooling to room temperature, the aqueous phase was separated off and the organic phase was concentrated by distilling out 60 ml of [MTB [sic]] MTB. The product was crystallized by adding 500 ml of heptane and slowly cooling to room temperature. The coarsely crystalline solid was filtered off with suction, washed with heptane and dried to constant weight in a vacuum oven at 40° C.

Yield: 108.9 g (80%), HPLC >99.5% area.

## Example 10

## S-2-Hydroxy-3-methoxy-3,3-diphenylpropionic acid (racemate resolution with L-proline methyl ester)

148.8 g of a 30% strength methanolic sodium methanolate solution (0.826 mol) were added dropwise to 240 g of a 57% strength methanolic L-proline methyl ester hydrochloride solution (0.826 mol) at room temperature, and 2.41 of MTB and 225 g (0.826 mol) of 2-hydroxy-3-methoxy-3,3-diphenylpropionic acid were added. After 2680 ml of MTB/methanol mixture had been distilled out with simultaneous dropwise addition of 2.4 l of MTB, the mixture was slowly cooled to room temperature, the crystals (R-2-hydroxy-3-methoxy-3,3-diphenylpropionic acid x L-proline methyl (ester) were filtered off with suction, and the solid was washed with 150 ml of MTB. The filtrate was concentrated by distilling out 1.5 l of MTB, and 1.0 l of water was added. The pH was adjusted

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to 1.2 with concentrated hydrochloric acid at room temperature and, after stirring and phase separation, the aqueous phase was separated off and extracted with 0.4 l of MTB. The combined organic phases were extracted with 0.4 l of water. The residue after the MTB had been stripped off was dissolved in 650 ml of toluene under reflux, and the product was crystallized by seeding and slow cooling. Filtration with suction, washing with toluene and drying in a vacuum oven resulted in 78.7 g of S-2-hydroxy-3-methoxy-3,3-diphenylpropionic acid (yield 35% based on the racemate).

Chiral HPLC: 100% pure; HPLC: 99.8%

## Example 11

S-2-Hydroxy-3-methoxy-3,3-diphenylpropionic acid (racemate resolution with (S)-1-(4-nitrophenyl)ethylamine)

30.5 g (0.184 mol) of (S)-1-(4-nitrophenyl)ethylamine were added to 100 g (0.368 mol) of 2-hydroxy-3-methoxy-3,3-diphenylpropionic acid in 750 ml of acetone and 750 ml of MTB under reflux, the mixture was seeded, boiled under reflux for one hour and slowly cooled to room temperature for crystallization. The crystals (S-2-hydroxy-3-methoxy-3,3-diphenylpropionic acid x (S)-1-(4-nitrophenyl)ethylamine) were filtered off with suction and washed with MTB. The residue was suspended in 500 ml of water and 350 ml of MTB and then the pH was adjusted to 1.2 with concentrated hydrochloric acid at room temperature, and, after stirring and phase separation, the aqueous phase was separated off and extracted with 150 ml of MTB. The combined organic phases were extracted with 100 ml of water. 370 ml of MTB were distilled out and then 390 ml of n-heptane were added under reflux, and the mixture was slowly cooled to room temperature while the product crystallized. Filtration with suction, washing with n-heptane and drying in a vacuum oven resulted in 35.0 g of S-2-hydroxy-3-methoxy-3,3-diphenylpropionic acid (yield 35% based on the racemate).

Chiral HPLC: 100% pure; HPLC: 99.8%

## Example 12

Benzyl 3-methoxy-2-(4-methoxy-6,7-dihydro-5H-cyclopentapyrimidin-2-yloxy)-3,3-diphenylpropionate

24.48 g (90 mmol) of 3-methoxy-3,3-diphenyl-2-hydroxypropionic acid were dissolved in 150 ml of DMF, and 13.7 g (99 mmol) of potassium carbonate were added. The suspension was stirred at room temperature for 30 min. Then 10.7 ml (90 mmol) of benzyl bromide were added dropwise over the course of 5 min, and the mixture was stirred for 1 h, during which the temperature rose to 32°C.

To this mixture were successively added 24.84 g (180 mmol) of K<sub>2</sub>CO<sub>3</sub> and 20.52 g (90 mmol) of 2-methanesulfonyl-4-methoxy-6,7-dihydro-5H-cyclopentapyrimidine [[sic]], and the mixture was stirred at 80°C. for 3 h.

For workup, the contents of the flask were diluted with about 600 ml of H<sub>2</sub>O and cautiously acidified with concentrated HCl, and 250 ml of ethyl acetate were added. 31.4 g of pure product precipitated and were filtered off.

The ethyl acetate phase was separated from the mother liquor, the aqueous phase was extracted again with ethyl acetate, and the combined organic phases were concentrated. The oily residue (19 g) was purified by chromatography (cyclohexane/ethyl acetate=9/1) to result in a further 10.5 g of pure product.

Total yield: 41.9 g (82.2 mmol)=91%; Melting point 143-147°C.; MS: MH<sup>+</sup>=511

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## Example 13

3-Methoxy-2-(4-methoxy-(6,7-dihydro-5H-cyclopentapyrimidin-2-yl-oxy)-3,3-diphenylpropionic acid

5 40 g (78.4 mmol) of benzyl 3-methoxy-2-(4-methoxy-6,7-dihydro-5H-cyclopentapyrimidin-2-yloxy)-3,3-diphenylpropionate were dissolved in 400 ml of ethyl acetate/methanol (4:1), about 500 mg of palladium on active carbon (10%) were added, and the mixture was exposed to a hydrogen atmosphere until no further gas was taken up. The catalyst was filtered off, the solution was evaporated, and the residue was crystallized from ether.

## Example 14

Ethyl 2S-3,3-diphenyloxirane-2-carboxylate

20 2.57 g (10.2 mmol) of ethyl 3,3-diphenylacrylate and 464 mg of 4-phenylpyridine N-oxide were dissolved in 24 ml of methylene chloride, and 432 mg (6.5 mol %) of (5,5)-(+)N,N'-bis(3,5-ditert-butylsalicylidene)-1,2-cyclohexanediaminomanganesc(III) chloride were added. While cooling in ice, 6.4 ml of a 12% strength sodium hypochlorite [[sic]] solution were added, and the mixture was stirred while cooling in ice for 30 min and at room temperature overnight. The solution was diluted to 200 ml with water, extracted with ether, dried and evaporated. 2.85 g of a colorless oil were obtained. Purification by [NPLC [sic]] HPLC (cyclohexane:ethyl acetate=9:1) resulted in 1.12 g of oil with an enantiomer ratio of about 8:1 in favor of the S configuration.

30 <sup>1</sup>H-NMR [CDCl<sub>3</sub>], δ=1.0 (t, 3H); 3.9 (m, 3H); 7.3 (m, 10H)

## Example 15

2-Methylsulfonyl-6,7-dihydro-5H-cyclopentapyrimidin-4-ol [[sic]]

35 46.9 g (330 mmol) of methyl cyclopentanone-2-carboxylate and 53.5 g (192 mmol) of 5-methylisothiourea [[sic]] sulfate were successively added to 29.6 g (528 mmol) of KOH in 396 ml of methanol, and the mixture was stirred at room temperature overnight, acidified with 1N hydrochloric acid and diluted with water. The crystals which separated out were filtered off with suction and dried. 20 g of crystals were obtained.

## Example 16

[[sulfanyl] Sulfanyl 4-[Chloro] chloro-2-methyl-6,7-dihydro-5H-cyclopentapyrimidine [[sic]]]

50 255 ml of phosphorus oxychloride were added to 20 g (110 mmol) [[lacuna]], and the mixture was stirred at 80°C. for 3 hours. Phosphorus oxychloride was evaporated off, ice was added to the residue, and the crystals which separated out were filtered off with suction. 18.5 g of a brownish solid were obtained.

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## Example 17

4-Methoxy-2-methylsulfonyl-6,7-dihydro-5H-cyclopentapyrimidine [[sic]]

60 18.05 g (90 mmol) of 4-chloro-2-methylsulfonyl-6,7-dihydro-5H-cyclopentapyrimidine [[sic]] were dissolved in 200 ml of methanol. At 45°C., 16.7 g of sodium methoxide (as 30% strength solutions [[sic]] in methanol) were added dropwise, and the mixture was stirred for 2 hours. The solution was evaporated, taken up in ethyl acetate and acidified with dilute hydrochloric acid, and the ethyl acetate extract was evaporated. 15.5 g of an oil remained.

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<sup>1</sup>H-NMR [DMSO],  $\delta$ =2.1 (quintet, 2H); 2.5 (s, 3H); 2.8 (dt, 4H); 3.9 (s, 3H) ppm

## Example 18

## 2-Methylsulfonyl-4-methoxy-6,7-dihydro-5H-cyclopento-pyrimidine [[sic]]

15 g (76.2 mmol) of 4-methoxy-2-methylsulfonyl-6,7-dihydro-5H-cyclopentapyrimidine [[sic]] were dissolved in 160 ml of glacial acetic acid/methylene chloride (1:1), and 1.3 g of sodium tungstate were added. At 35°C., 17.5 ml (170 ml [[sic]]) of a 30% strength  $H_2O_2$  solution were added dropwise. The mixture was then diluted with 500 ml of water and 100 ml of methylene chloride, and the organic phase was separated off, dried and evaporated. 14 g of oil remained and were crystallized from ether.

<sup>1</sup>H-NMR [CDCl<sub>3</sub>],  $\delta$ =2.2 (quintet, 2H); 3.0 (dt., 4H); 3.3 (s, 3H); 4.1 (s, 3H) ppm

## Example 19

## 1-Benzene sulfonyl-3-(4,6-dimethoxy-2-pyrimidinyloxy)-4-methoxy-4,4-diphenyl-2-butanone

0.37 g (2.4 mmol) of phenyl methane [[sic]] sulfone were dissolved in 10 ml of dry THF and then, at -70°C., 2 eq. of butyllithium (2.94 ml; 1.6 molar solution in hexane) were added dropwise. After 1 h at -70°C., 1 g (2.4 mmol) of methyl 2-(4,6-dimethoxy-2-pyrimidinyloxy)-3-methoxy-3,3-diphenylpropionate [[sic]] dissolved in 5 ml of THF was added dropwise. The reaction mixture was then stirred at -70°C. for 1 h and at -10°C. for 1 h and then warmed to room temperature. For workup, about 10 ml of saturated NH<sub>4</sub>Cl solution were added dropwise, thorough extraction with ethyl acetate was carried out, and the combined organic phases [[lacuna]] with-saturated N-Cl [[sic]] solution and dried over Na<sub>2</sub>SO<sub>4</sub>. The residue obtained after drying and concentration was purified by chromatography on silica gel (n-heptane/ethyl acetate 15%→30%) and subsequently [MPLC] HPLC on RP silica gel (acetonitrile/H<sub>2</sub>O+TFA); 0.3 g of a white amorphous powder was obtained as product.

## Example 20

## 3,3-Diphenyloxiram-2-carbonitrile [[sic]]

3.1 g (54.9 mmol) of sodium methoxide were suspended in 20 ml of dry THF and then, at -10°C., a mixture of 5 g (27.4 mmol) of benzophenone and 4.2 g (54.9 mmol) of chloroacetonitrile was added dropwise.

The reaction mixture was stirred at -10°C. for about 2 h, then poured into water and extracted several times with ethyl acetate. The combined organic phases were dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated, and the residue was purified by chromatography on silica gel (n-heptane/ethyl acetate).

Yield: 1.2 g (20%)

<sup>1</sup>H-NMR [CDCl<sub>3</sub>],  $\delta$ =3.9 (s, 1H); 7.4-7.5 (m, 10 H) ppm

## Example 21

## 2-Hydroxy-3-methoxy-3,3-diphenylpropionitrile

6.5 [[lacuna]] g (29.4 mmol) of 3,3-diphenyloxiran-2-carbonitrile were dissolved in 60 ml of methanol and, at 0°C., about 2 ml of boron trifluoride etherate solution were added. The mixture was stirred further at 0°C. for 1 h and then at room temperature overnight. For workup it was diluted with diethyl ether and washed with saturated NaCl solution, and the organic phase was dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated.

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The residue comprised 7.3 g of a white amorphous powder which was used directly in the subsequent reactions.

<sup>1</sup>H-NMR [[CDCl<sub>3</sub>]] / [CDCl<sub>3</sub>],  $\delta$ =2.95 (broad s, OH), 3.15 (s, 3H), 5.3 (s, 1H), 7.3-7.5 (m, 10) ppm

## Example 22

## 2-(4,6-Dimethoxy-2-pyrimidinyloxy)-3-methoxy-3,3-diphenylpropionitrile

7.3 g (28.8 mmol) of 2-hydroxy-3-methoxy-3,3-diphenylpropionitrile were dissolved in 90 ml of DMF, and 4 g (28.8 mmol) of K<sub>2</sub>CO<sub>3</sub> and 6.3 g (28 mmol) of 2-methanesulfonyl-4,6-dimethoxypyrimidine were added. The mixture was stirred at room temperature for about 12 h, then poured into water and extracted with ethyl acetate. The combined organic phases were washed again with H<sub>2</sub>O, dried and concentrated. The residue obtained in this way was then purified by chromatography on silica gel (n-heptane/ethyl acetate).

Yield: 6.9 g of white amorphous powder

FAB-MS: 392 (M+H<sup>+</sup>) <sup>1</sup>H-NMR [CDCl<sub>3</sub>],  $\delta$ =3.3 (s, 3H); 4.95 (s, 6H), 5.85 (s, 1H); 6.3 (s, 1H); 7.3-7.5 (m, 10H) ppm

## Example 23

## 5-[2-(4,6-Dimethoxy-2-pyrimidinyloxy)-3-methoxy-3,3-diphenylpropyl]-1H-tetrazole [[sic]]

0.5 g (1.3 mmol) of nitrile was dissolved in 10 ml of toluene, and 85 mg (1.3 mmol) of NaN<sub>3</sub> and 460 mg (1.4 mmol) of Bu<sub>3</sub>SnCl were successively added, and then the mixture was refluxed for about 40 h. Cooling was followed by dilution with ethyl acetate and washing with 10% aqueous KF solution and with NaCl solution. After drying over MgSO<sub>4</sub> and concentration there remained 1.0 g of a yellow oil, which was purified by chromatography on silica gel (n-heptane/ethyl acetate).

Concentration of the fractions resulted in 60 mg of the 1H-tetrazole and 110 mg of the 1-methyltetrazole, each as amorphous white solids.

## 5-[2-(4,6-Dimethoxy-2-pyrimidinyloxy)-3-methoxy-3,3-diphenylpropyl]-1H-tetrazole [[sic]]

Electrospray-MS: 435 (M+H<sup>+</sup>) <sup>1</sup>H-NMR (CDCl<sub>3</sub>):  $\delta$  (ppm) 3.28 (s, 3H), 3.85 (s, 6H), 5.75 (s, 1H), 7.25-7.40 (m, 10H), 7.50 (s, 1H).

## 5-[2-(4,6-Dimethoxy-2-pyrimidinyloxy)-3-methoxy-3,3-diphenylpropyl]-1-methyltetrazole [[sic]]

Electrospray-MS: 471 (M+H<sup>+</sup>) <sup>1</sup>H-NMR (CDCl<sub>3</sub>):  $\delta$  (ppm) 3.0 (s, 3H), 3.35 (s, 3H) [[sic]], 3.80 (s, 6H), 5.75 (s, 1H), 7.30-7.40 (m, 11H).

## Example 24

## 2-(4,6-Dimethoxy-2-pyrimidinyloxy)-3-methylsulfinyl-3,3-diphenylpropionic acid

1.2 g (2.9 mmol) of 2-(4,6-dimethoxy-2-pyrimidinyloxy)-3-methylsulfonyl-3,3-diphenylpropionic [[sic]] acid were introduced into 15 ml of glacial acetic acid at 0°C. and 294  $\mu$ l of 30% strength  $H_2O_2$  were added dropwise. The mixture was stirred at room temperature overnight, poured into water, extracted with CH<sub>2</sub>Cl<sub>2</sub> and washed with sodium thiosulfate solution and brine. After drying, 1 g of substance was isolated as a white foam.

2-(4,6-Dimethoxy-2-pyrimidinyloxy)-3-methylsulfonyl-3,3-diphenylpropionic acid

0.6 g (1.45 mmol) of 2-(4,6-dimethoxy-2-pyrimidinyloxy)-3-methylsulfonyl-3,3-diphenylpropionic **[[sic]]** acid was introduced into 15 ml of glacial acetic acid at room

temperature, and 294  $\mu$ l of 30% strength  $H_2O_2$  were added dropwise. The mixture was stirred at room temperature overnight, heated at 50° C. for a further 3 h, poured into water and washed with sodium thiosulfate solution and brine. After drying, 400 mg were isolated as a white solid.

The compounds listed in Table 1 **[[sic]]** can be prepared in a similar way.

TABLE 1

No.	$R^1$	$R^4, R^5$	$R^6$	$R^2$	$R^3$	$X$	$Y$	$Z$	m.p. [°C.]
<b>I-195</b> <b>I-1</b>	$[\text{OMe}]OCH_3$	Phenyl	Methyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	81
<b>I-196</b> <b>I-2</b>	OH	Phenyl	Methyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	167
<b>I-197</b> <b>I-3</b>	OH	Phenyl	$\text{CH}_2\text{—CH}_2\text{—S—CH}_3$	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-198</b> <b>I-4</b>	OH	Phenyl	Ethyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	81 (decomp.)
<b>I-199</b> <b>I-5</b>	OH	Phenyl	iso-Propyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	182
<b>I-200</b> <b>I-6</b>	OH	Phenyl	Methyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	S	168
<b>I-201</b> <b>I-7</b>	OH	Phenyl	$\text{CH}_2\text{—CH}_2\text{—SO}_2\text{—}$	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-202</b> <b>I-8</b>	OH	Phenyl	$\text{CH}_2\text{—CH}_2\text{—SO}_2\text{—}$	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	S	O	
<b>I-203</b> <b>I-9</b>	OH	Phenyl	$\text{CH}_2\text{—CH}_2\text{—SO}_2\text{—}$	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	$\text{C—CH}(\text{CH}_3)_2$	O	O	
<b>I-204</b> <b>I-10</b>	OH	Phenyl	$\text{CH}_2\text{—CH}_2\text{—SO}_2\text{—}$	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	$\text{C—CH}(\text{CH}_3)_2$	O	O	
<b>I-205</b> <b>I-11</b>	OH	Phenyl	$\text{CH}_2\text{—CH}_2\text{—SO}_2\text{—}$	$[\text{OMe}]OCH_3$	$\text{NH}\bullet\text{OCH}_3$	CH	O	O	
<b>I-206</b> <b>I-12</b>	OH	Phenyl	n-Propyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	174
<b>I-207</b> <b>I-13</b>	$[\text{OMe}]OCH_3$	Phenyl	n-Propyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-208</b> <b>I-14</b>	OH	Phenyl	n-Propyl	$[\text{OEt}]OC_2H_5$	$[\text{OEt}]OC_2H_5$	CH	O	O	
<b>I-209</b> <b>I-15</b>	OH	Phenyl	n-Butyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-210</b> <b>I-16</b>	OH	Phenyl	iso-Butyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-211</b> <b>I-17</b>	OH	Phenyl	iso-Butyl	$[\text{OMe}]OCH_3$	$\text{O—CH}_2\text{—CH}_2\text{—C}$		O	O	
<b>I-212</b> <b>I-18</b>	OH	Phenyl	tert.-Butyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-213</b> <b>I-19</b>	OH	Phenyl	Cyclopropyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-214</b> <b>I-20</b>	OH	Phenyl	Cyclopentyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-215</b> <b>I-21</b>	OH	Phenyl	Cyclohexyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-216</b> <b>I-22</b>	OH	Phenyl	$(\text{CH}_3)_3\text{C—CH}_2\text{—CH}_2$	$[\text{OEt}]OC_2H_5$	$[\text{OEt}]OC_2H_5$	CH	O	O	
<b>I-217</b> <b>I-23</b>	OH	Phenyl	$(\text{CH}_3)_2\text{CH—CH}_2\text{—}$	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	173
<b>I-218</b> <b>I-24</b>	OH	Phenyl	$\text{HO—CH}_2\text{—CH}_2$	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-219</b> <b>I-25</b>	OH	Phenyl	$\text{HO}_2\text{C—CH}_2\text{—}$	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-220</b> <b>I-26</b>	OH	Phenyl	Cyclopropyl-methylene <b>[[sic]]</b>	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	115
<b>I-221</b> <b>I-27</b>	OH	Phenyl	H	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-222</b> <b>I-28</b>	OH	Phenyl	Methyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	—	
<b>I-223</b> <b>I-29</b>	OH	Phenyl	Phenyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	136
<b>I-224</b> <b>I-30</b>	OH	Phenyl	Phenyl	$[\text{OMe}]OCH_3$	$\text{O—CH}(\text{CH}_3)\text{—CH}_2\text{—C}$		O	O	
<b>I-225</b> <b>I-31</b>	OH	Phenyl	Phenyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-226</b> <b>I-32</b>	OH	Phenyl	4-Isopropyl-Phenyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-227</b> <b>I-33</b>	OH	Phenyl	4-Methyl-S-Phenyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-228</b> <b>I-34</b>	OH	Phenyl	4-Methyl-O-Phenyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-229</b> <b>I-35</b>	OH	Phenyl	3-Ethyl-Phenyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-230</b> <b>I-36</b>	OH	Phenyl	2-Methyl-Phenyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-231</b> <b>I-37</b>	OH	Phenyl	2-Cl-Phenyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-232</b> <b>I-38</b>	OH	Phenyl	3-Br-Phenyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-233</b> <b>I-39</b>	OH	Phenyl	4-F-Phenyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-234</b> <b>I-40</b>	OH	Phenyl	4-F-Phenyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	S	O	
<b>I-235</b> <b>I-41</b>	OH	Phenyl	4-CH <sub>3</sub> -Phenyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-236</b> <b>I-42</b>	OH	Phenyl	3-NO <sub>2</sub> -Phenyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-237</b> <b>I-43</b>	OH	Phenyl	2-HO-Phenyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-238</b> <b>I-44</b>	OH	Phenyl	3,4-Dimethoxyphenyl	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	
<b>I-239</b> <b>I-45</b>	OH	Phenyl	3,4-Dioxomethylene-phenyl- <b>[[sic]]</b>	$[\text{OMe}]OCH_3$	$[\text{OMe}]OCH_3$	CH	O	O	

TABLE 1-continued

[I]									
No.	R <sup>1</sup>	R <sup>4</sup> , R <sup>5</sup>	R <sup>6</sup>	R <sup>2</sup>	R <sup>3</sup>	X	Y	Z	m.p. [°C.]
[I-240]I-46	OH	Phenyl	3,4,5-Triimethoxyphenyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-241]I-47	OH	Phenyl	Benzyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-242]I-48	OH	Phenyl	2-Cl-Benzyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-243]I-49	OH	Phenyl	3-Br-Benzyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-244]I-50	OH	Phenyl	4-F-Benzyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-245]I-51	OH	Phenyl	2-Methyl-Benzyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-246]I-52	OH	Phenyl	2-Methyl-Benzyl	[OMe]OCH <sub>3</sub>	O—CH=CH—C		O	O	
[I-247]I-53	OH	Phenyl	3-Ethyl-Benzyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-248]I-54	OH	Phenyl	4-iso-Propyl-Benzyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-249]I-55	OH	Phenyl	4-NO <sub>2</sub> -Propyl-Benzyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-250]I-56	OH	Phenyl	2-Methyl-5-Propyl-Benzyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-251]I-57	OH	Phenyl	2-Methyl-5-Propyl-Benzyl	[OEt]OC <sub>2</sub> H <sub>5</sub>	[OEt]OC <sub>2</sub> H <sub>5</sub>	CH	O	O	
[I-252]I-58	OH	Phenyl	4-Methyl-2-Propyl-Benzyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-253]I-59	OH	Phenyl	3,4-Dioxomethyl-enebenzyl [[sic]]	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-254]I-60	OH	4-F-Phenyl	4-Methyl-2-Propyl-Benzyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	163-165 (decomp.)
[I-255]I-61	[OMe]OCH <sub>3</sub>	4-F-Phenyl	Methyl	[OEt]OC <sub>2</sub> H <sub>5</sub>	[OEt]OC <sub>2</sub> H <sub>5</sub>	CH	O	O	
[I-256]I-62	OH	4-Cl-Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-257]I-63	OH	4-Methyl—O—Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-258]I-64	OH	4-Methyl—O—Phenyl	Ethyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-259]I-65	OH	4-Methyl-Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-260]I-66	OH	4-Methyl-Phenyl	Methyl	[OMe]OCH <sub>3</sub>	O—CH <sub>2</sub> —CH <sub>2</sub> —C		O	O	
[I-261]I-67	OH	3-CF <sub>3</sub> -Phenyl	n-Propyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-262]I-68	OH	3-CF <sub>3</sub> -Phenyl	n-Propyl	[OMe]OCH <sub>3</sub>	O—CH(CH <sub>3</sub> )—CH <sub>2</sub> —C		O	O	
[I-263]I-69	OH	4-NO <sub>2</sub> -Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-264]I-70	OH	4-NO <sub>2</sub> -Phenyl	Methyl	[OMe]OCH <sub>3</sub>	O—CH=CH—C		O	O	
[I-265]I-71	OH	3-Cl-Phenyl	Ethyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-266]I-72	OH	2-F-Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	193-194 (decomp.)
[I-267]I-73	OH	2-F-Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	S	O	
[I-268]I-74	OH	2-Methyl—O—Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-269]I-75	OH	2-Methyl—O—Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	S	
[I-270]I-76	OH	3,4-Dimethoxyphenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-271]I-77	OH	3,4-Dioxomethyl-enephenyl [[sic]]	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-272]I-78	OH	p-CF <sub>3</sub> -Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-273]I-79	OH	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OEt]OC <sub>2</sub> H <sub>5</sub>	CH	O	O	
[I-274]I-80	[OMe]OCH <sub>3</sub>	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OEt]OC <sub>2</sub> H <sub>5</sub>	CH	O	O	
[I-275]I-81	OH	Phenyl	Ethyl	[OMe]OCH <sub>3</sub>	NH—[OMe]	CH	O	O	
[I-276]I-82	OH	p-Methyl—O—Phenyl	n-Propyl	[OMe]OCH <sub>3</sub>	OCF <sub>3</sub>	CH	O	O	
[I-277]I-83	OH	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	CF <sub>3</sub>	CH	O	O	
[I-278]I-84	OH	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	CF <sub>3</sub>	N	O	O	
[I-279]I-85	OH	3,4-Dimethoxyphenyl	Benzyl	Methyl	Methyl		O	O	
[I-280]I-86	OH	3,4-Dimethoxyphenyl	Methyl	[OMe]OCH <sub>3</sub>	O—CH <sub>2</sub> CH <sub>2</sub> —C		O	O	
[I-281]I-87	OH	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	O—CH <sub>2</sub> —CH <sub>2</sub> —C		O	O	126 (decomp.)
[I-282]I-88	OH	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	O—CH(CH <sub>3</sub> )—CH <sub>2</sub> —C		O	O	
[I-283]I-89	OH	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	N(CH <sub>3</sub> )—CH=CH—C		O	O	118
[I-284]I-90	OH	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	S—C(CH <sub>3</sub> )=C(CH <sub>3</sub> )—C		O	O	
[I-285]I-91	OH	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	O—C(CH <sub>3</sub> )=CH—C		O	O	
[I-286]I-92	OH	Phenyl	Methyl	Methyl	O—C(CH <sub>3</sub> )=CH—C		O	O	
[I-287]I-93	OH	Phenyl	Methyl	Methyl	O—CH=CH—C		O	O	
[I-288]I-94	OH	4-F-Phenyl	Methyl	Methyl	S—CH=CH—C		O	O	
[I-289]I-95	OH	4-F-Phenyl	H	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	S	
[I-290]I-96	OH	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	CH <sub>2</sub> —CH <sub>2</sub> —CH <sub>2</sub> —C		O	O	149-151 (decomp.)
[I-291]I-97	OH	Phenyl	Methyl	Methyl	CH <sub>2</sub> —CH <sub>2</sub> —CH <sub>2</sub> —C		O	O	157 (decomp.)
[I-292]I-98	OH	Phenyl	Methyl	Ethyl	CH <sub>2</sub> —CH <sub>2</sub> —CH <sub>2</sub> —CH <sub>2</sub> —C		O	O	

TABLE 1-continued

I									
No.	R <sup>1</sup>	R <sup>4</sup> , R <sup>5</sup>	R <sup>6</sup>	R <sup>2</sup>	R <sup>3</sup>	X	Y	Z	m.p. [°C.]
I-293	J-99 OH	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	CH <sub>2</sub> —CH <sub>2</sub> —CH <sub>2</sub> —CH <sub>2</sub> —C	O	O	O	
I-294	J-100 OH	Phenyl	Methyl	Methyl	CH	O	O	O	
I-295	J-101 OH	Phenyl	Methyl	Ethyl	CH	O	O	O	
I-296	J-102 OH	Phenyl	Methyl	Methyl	Methyl	O	O	O	
I-297	J-103 OH	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[Me]CH <sub>3</sub>	CH	O	O	
I-298	J-104 OH	Cyclohexyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
I-299	J-105 OH	Cyclohexyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH <sub>2</sub> —CH <sub>2</sub> —CH <sub>2</sub> —C	O	O	
I-300	J-106 OH	Phenyl	Methyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	S	S	
I-301	J-107 OH	Phenyl	Methyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	O	S	134
I-302	J-108 OCH <sub>3</sub>	Phenyl	Methyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	S	S	
I-303	J-109 OH	Phenyl	Methyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	O	O	
I-304	J-110 OCH <sub>3</sub>	2-Fluorophenyl	Methyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	O	O	
I-305	J-111 OC <sub>2</sub> H <sub>5</sub>	3-Chlorophenyl	Methyl	OCH <sub>3</sub>	OCH <sub>3</sub>	N	O	O	
I-306	J-112 ON(CH <sub>3</sub> ) <sub>2</sub>	4-Bromophenyl	Methyl	CF <sub>3</sub>	CF <sub>3</sub>	CH	S	O	
I-307	J-113 O—CH <sub>2</sub> —C≡CH	Phenyl	Ethyl	OCH <sub>3</sub>	CF <sub>3</sub>	CH	O	O	
I-308	J-114 OH	Phenyl	Propyl	OCH <sub>3</sub>	OCF <sub>3</sub>	CH	O	S	
I-309	J-115 OCH <sub>3</sub>	Phenyl	i-Propyl	OCH <sub>3</sub>	CH <sub>3</sub>	CH	O	O	
I-310	J-116 OC <sub>2</sub> H <sub>5</sub>	Phenyl	s-Butyl	OCH <sub>3</sub>	Cl	CH	S	O	
I-311	J-117 ON(CH <sub>3</sub> ) <sub>2</sub>	2-Methylphenyl	Methyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	O	O	
I-312	J-118 ON(CH <sub>3</sub> ) <sub>2</sub>	3-Methoxyphenyl	Methyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	O	O	
I-313	J-119 ON—C(CH <sub>3</sub> ) <sub>2</sub>	4-Nitrophenyl	Methyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	O	O	
I-314	J-120 ON(CH <sub>3</sub> ) <sub>2</sub>	Phenyl	1-Phenylpropyn-3-yl	OCH <sub>3</sub>	OCF <sub>3</sub>	N	O	S	
I-315	J-121 ON—C(CH <sub>3</sub> ) <sub>2</sub>	2-Hydroxyphenyl	Methyl	OCH <sub>3</sub>	CH <sub>3</sub>	N	O	O	
I-316	J-122 ONSO <sub>2</sub> (C <sub>6</sub> H <sub>5</sub> ) <sub>2</sub>	3-Trifluoromethyl-phenyl	Methyl	OCH <sub>3</sub>	Cl	N	O	O	
I-317	J-123 NH[Phenyl]—phenyl	4-Dimethyl-aminophenyl	Methyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	S	O	
I-318	J-124 OC <sub>2</sub> H <sub>5</sub>	Phenyl	Trifluoromethyl	CH <sub>3</sub>	CH <sub>3</sub>	CH	O	O	
I-319	J-125 ON(CH <sub>3</sub> ) <sub>2</sub>	Phenyl	Benzyl	Cl	Cl	CH	O	O	
I-320	J-126 ON(CH <sub>3</sub> ) <sub>2</sub>	Phenyl	2-Methoxyethyl	OCH <sub>3</sub>	—O—CH <sub>2</sub> —CH <sub>2</sub> —	S	O		
I-321	J-127 OH	Phenyl	Phenyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	O	O	
I-322	J-128 OH	Phenyl	Phenyl	OCH <sub>3</sub>	—O—CH <sub>2</sub> —CH <sub>2</sub> —	O	O		
I-323	J-129 OH	Phenyl	Phenyl	OCH <sub>3</sub>	OCH <sub>3</sub>	N	O	O	
I-324	J-130 OH	Phenyl	Phenyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	S	O	
I-325	J-131 OH	Phenyl	Phenyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	S	S	
I-326	J-132 OH	Phenyl	Phenyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	O	S	
I-327	J-133 OH	Phenyl	Phenyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	O	O	
I-328	J-134 OH	Phenyl	Phenyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	O	O	
I-329	J-135 OH	—(CH <sub>2</sub> ) <sub>5</sub> —	Phenyl	Phenyl	OCH <sub>3</sub>	CH	O	O	
I-330	J-136 OH	Phenyl	[2-thiophenyl]—2-thiazolyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	O	O	
I-331	J-137 OCH <sub>3</sub>	2-Fluorophenyl	Phenyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	O	O	
I-332	J-138 OC <sub>2</sub> H <sub>5</sub>	3-Chlorophenyl	Phenyl	OCH <sub>3</sub>	OCH <sub>3</sub>	N	O	O	
I-333	J-139 ON(CH <sub>3</sub> ) <sub>2</sub>	4-Bromophenyl	Phenyl	CF <sub>3</sub>	CF <sub>3</sub>	CH	O	O	
I-334	J-140 [OCH <sub>2</sub> CH] OCH <sub>2</sub> =CH	Phenyl	2-Fluorophenyl	OCH <sub>3</sub>	CF <sub>3</sub>	CH	O	O	
I-335	J-141 OH	Phenyl	3-Chlorophenyl	OCH <sub>3</sub>	OCF <sub>3</sub>	CH	O	S	
I-336	J-142 OCH <sub>3</sub>	Phenyl	4-Bromophenyl	OCH <sub>3</sub>	CH <sub>3</sub>	CH	O	O	
I-337	J-143 OC <sub>2</sub> H <sub>5</sub>	Phenyl	4-Thiazolyl	OCH <sub>3</sub>	Cl	CH	S	O	
I-338	J-144 ON(CH <sub>3</sub> ) <sub>2</sub>	2-Methylphenyl	Phenyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	O	O	
I-339	J-145 ON—C(CH <sub>3</sub> ) <sub>2</sub>	2-Methoxyphenyl	Phenyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	O	O	
I-340	J-146 OH	Phenyl	Methyl	OCH <sub>3</sub>	—CH <sub>2</sub> —CH <sub>2</sub> —CH <sub>2</sub> —C	O	O		
I-341	J-147 OH	4-Fluorophenyl	Methyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	O	O	168 (decomp.)
I-342	J-148 OH	4-Fluorophenyl	Methyl	OCH <sub>3</sub>	—CH <sub>2</sub> —CH <sub>2</sub> —CH <sub>2</sub> —C	O	O		
I-343	J-149 NH—SO—C <sub>6</sub> H <sub>5</sub>	4-Nitrophenyl	Phenyl	OCH <sub>3</sub>	CH	O	O		
I-344	J-150 OCH <sub>3</sub>	Phenyl	3-Imidazolyl	OCH <sub>3</sub>	—O—CH <sub>2</sub> —CH <sub>2</sub> —	O	O		
I-345	J-151 OC <sub>2</sub> H <sub>5</sub>	Phenyl	4-Imidazolyl	OCH <sub>3</sub>	CF <sub>3</sub>	N	S	O	
I-346	J-152 ON(CH <sub>3</sub> ) <sub>2</sub>	Phenyl	2-Pyrazolyl	OCH <sub>3</sub>	OCF <sub>3</sub>	N	O	S	
I-347	J-153 ON—C(CH <sub>3</sub> ) <sub>2</sub>	2-Hydroxyphenyl	Phenyl	OCH <sub>3</sub>	CH <sub>3</sub>	N	O	O	
I-348	J-154 NH—SO <sub>2</sub> —C <sub>6</sub> H <sub>5</sub>	3-Trifluoromethyl-phenyl	Phenyl	OCH <sub>3</sub>	Cl	N	O	O	
I-349	J-155 NH[Phenyl]—phenyl	4-Dimethylamino-phenyl	Phenyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	S	O	
I-350	J-156 ONa	Phenyl	Phenyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	S	S	

TABLE 1-continued

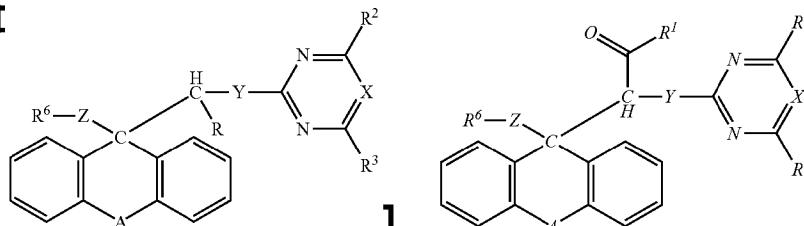
[

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No.	R <sup>1</sup>	R <sup>4</sup> , R <sup>5</sup>	R <sup>6</sup>	R <sup>2</sup>	R <sup>3</sup>	X	Y	Z	m.p. [°C.]
[I-351]I-157	O—CH <sub>2</sub> —C≡C	Phenyl	Phenyl	OCH <sub>3</sub>	OCH <sub>3</sub>	N	S	S	
[I-352]I-158	OH	Phenyl	Phenyl	CF <sub>3</sub>	CF <sub>3</sub>	CH	O	S	
[I-353]I-159	OCH <sub>3</sub>	Phenyl	Phenyl	OCF <sub>3</sub>	OCF <sub>3</sub>	CH	O	O	
[I-354]I-160	OC <sub>2</sub> H <sub>5</sub>	Phenyl	2-Dimethylamino-phenyl	CH <sub>3</sub>	CH <sub>3</sub>	CH	O	O	
[I-355]I-161	ONC(CH <sub>3</sub> ) <sub>2</sub>	Phenyl	3-Hydroxyphenyl	Cl	Cl	CH	O	O	
[I-356]I-162	ON=C(CH <sub>3</sub> ) <sub>2</sub>	Phenyl	4-Trifluoromethyl-phenyl	OCH <sub>3</sub>	—O—CH <sub>2</sub> —CH <sub>2</sub> —		S	O	
[I-357]I-163	NH—SO <sub>2</sub> —C <sub>6</sub> H <sub>5</sub>	Phenyl	2-Oxazolyl	OCH <sub>3</sub>	CF <sub>3</sub>	N	S	S	
[I-358]I-164	OH	Phenyl	Methyl	CH <sub>3</sub>	CH <sub>3</sub>	CH	O	O	
[I-359]I-165	OH	Cyclohexyl	Methyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	O	O	
[I-360]I-166	OH	Cyclohexyl	Methyl	OCH <sub>3</sub>	CH <sub>2</sub> —CH <sub>2</sub> —CH—C		O	O	
[I-361]I-167	OH	Phenyl	Methyl	N(CH <sub>3</sub> ) <sub>2</sub>	N(CH <sub>3</sub> ) <sub>2</sub>	CH	O	O	
[I-362]I-168	OH	Phenyl	Methyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	O	SO <sub>2</sub>	
[I-363]I-169	OH	Phenyl	Methyl	OCH <sub>3</sub>	OCH <sub>3</sub>	CH	O	SO <sub>2</sub>	
[I-364]I-170	OH	3-F—Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-365]I-171	OH	3-F—Phenyl	Methyl	[OMe]OCH <sub>3</sub>	CH <sub>2</sub> —CH <sub>2</sub> —CH <sub>2</sub> —C		O	O	
[I-366]I-172	OH	4-F—Phenyl	Methyl	[OMe]OCH <sub>3</sub>	CH <sub>2</sub> —CH <sub>2</sub> CH <sub>2</sub> —C		O	O	142-143
									191°C.
[I-367]I-173	OH	3-Methyl—O—Phenyl	Methyl	[OMe]OCH <sub>3</sub>	CH <sub>2</sub> —CH <sub>2</sub> —CH <sub>2</sub> —C		O	O	158-161
									(decomp.)
[I-368]I-174	OH	3-Methyl—O—Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-369]I-175	OH	3-Methyl—O—Phenyl	Ethyl	[OMe]OCH <sub>3</sub>	CH <sub>2</sub> —CH <sub>2</sub> —CH <sub>2</sub> —C		O	O	
[I-370]I-176	OH	Phenyl	HO—CH <sub>2</sub> —CH <sub>2</sub>	[OMe]OCH <sub>3</sub>	CH <sub>2</sub> —CH <sub>2</sub> —CH <sub>2</sub> —C		O	O	
[I-371]I-177	OH	Phenyl	Methyl	[NMe <sub>2</sub> ]	[NMe <sub>2</sub> ]	N	O	O	181
				N(CH <sub>3</sub> ) <sub>2</sub>	N(CH <sub>3</sub> ) <sub>2</sub>				
[I-372]I-178	OH	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	N	O	O	
[I-323]I-179	OH	3-F—Phenyl	Methyl	OCH <sub>3</sub>	CH <sub>3</sub>	CH	O	O	
[I-374]I-180	NH—SO <sub>2</sub> — Phenyl	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CII	O	O	
[I-375]I-181	NH—SO <sub>2</sub> —[Me] CH <sub>3</sub>	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-376]I-182	CH <sub>2</sub> —SO <sub>2</sub> — Phenyl	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-377]I-183	NH—SO <sub>2</sub> —[Me] CH <sub>3</sub>	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-378]I-184	—CN	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-379]I-185	Tetrazole[[sic]]	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-380]I-186	NH—SO <sub>2</sub> — Phenyl	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	167
[I-381]I-187	N—Methyltetra- azole[[sic]]	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	
[I-382]I-188	ONa	Phenyl	Methyl	[OMe]OCH <sub>3</sub>	—O—CH <sub>2</sub> —CH <sub>2</sub> —C—		O	O	122-139
									(decomp.)
									[(zers.)]
[I-383]I-189	OH	o-F—Phenyl	Methyl	[OMe]OCH <sub>3</sub>	—O—CH <sub>2</sub> —CH <sub>2</sub> —C—		O	O	140-144
									(decomp.)
[I-384]I-190	OH	m-Methyl-Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[OMe]OCH <sub>3</sub>	CH	O	O	169-177
[I-385]I-191	OH	m-Methyl-Phenyl	Methyl	[OMe]OCH <sub>3</sub>	—O—CH <sub>2</sub> —CH <sub>2</sub> —C—		O	O	119-135
									(decomp.)
[I-386]I-192	OH	p-F—Phenyl	Methyl	[OMe]OCH <sub>3</sub>	[Me]CH <sub>3</sub>	CH	O	O	137-140
									(decomp.)
[I-387]I-193	OH	m-F—Phenyl	Methyl	Methyl	—O—CH <sub>2</sub> —CH <sub>2</sub> —C—		O	O	150-152
[I-388]I-194	OH	p-F—Phenyl	Methyl	Methyl	—O—CH <sub>2</sub> —CH <sub>2</sub> —C—		O	O	169-170

TABLE II

[



No.	R <sup>1</sup>	A	R <sup>6</sup>	R <sup>2</sup>	R <sup>3</sup>	X	Y	Z	m.p. [°C.]
II-1	OH	Bond	Methyl	OMethyl	OMethyl	CH	O	O	96-98
II-2	OH	CH <sub>2</sub>	Methyl	OMethyl	OMethyl	CH	O	O	
II-3	OH	CH <sub>2</sub> —CH <sub>2</sub>	Methyl	OMethyl	OMethyl	CH	O	O	
II-4	OH	CH=CH	Methyl	OMethyl	OMethyl	CH	O	O	
II-5	OH	O	Methyl	OMethyl	OMethyl	CH	O	O	
II-6	OH	S	Methyl	OMethyl	OMethyl	CH	O	O	
II-7	OH	NH(CH <sub>3</sub> )	Methyl	OMethyl	OMethyl	CH	O	O	
II-8	OH	Bond	Isopropyl	OMethyl	OMethyl	CH	O	O	137-139
II-9	OH	Bond	p-Isopropylphenyl	OMethyl	OMethyl	CH	O	O	
II-10	OH	Bond	Benzyl	OMethyl	OMethyl	CH	O	O	
II-11	OH	CH=CH	Ethyl	OMethyl	OMethyl	CH	O	O	
II-12	OH	CH=CH	(CH <sub>3</sub> ) <sub>2</sub> —CH <sub>2</sub> —CH <sub>2</sub>	OMethyl	OMethyl	CH	O	O	
II-13	OH	CH=CH	Cyclopropylmethyl [ene][sic]	OMethyl	OMethyl	CH	O	O	
II-14	OH	CH=CH	Methyl	OMethyl	O—CH <sub>2</sub> —CH <sub>2</sub> —C	O	O		
II-15	OH	CH <sub>2</sub> —CH <sub>2</sub>	Ethyl	OMethyl	O—CH=CH—C	O	O		
II-16	OH	[CH <sub>2</sub> =CH <sub>2</sub> ] CH <sub>2</sub> —CH <sub>2</sub>	Methyl	OMethyl	CH <sub>2</sub> —CH <sub>2</sub> —CH <sub>2</sub> —C	O	O		
II-17	OH	Bond	Methyl	OMethyl	CH <sub>2</sub> —CH <sub>2</sub> —CH <sub>2</sub> —C	O	O		147

Example 35

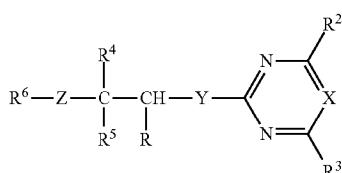
40 We claim:  
1. A compound of the formula I

Receptor binding data were measured by the binding assay described above for the compounds listed below. The results are shown in Table 2 [sic].

TABLE 2

Receptor binding data (K<sub>i</sub> values)

Compound	ET <sub>A</sub> [nM]	ET <sub>B</sub> [nM]
I-2	6	34
I-29	86	180
I-5	12	160
I-4	7	2500
I-87	1	57
I-89	86	9300
I-103	0.4	29
I-107	3	485
I-12	19	1700
I-26	23	2000
I-23	209	1100
I-47	150	1500
I-60	33	970
I-96	0.6	56
II-3	107	7300
II-1	28	2300



where

R is formyl, tetrazole, nitrile, [a COOH group]—CO<sub>2</sub>H or a radical which can be hydrolyzed to [COOH, and the other substituents have the following meanings: ]—CO<sub>2</sub>H;

R<sup>2</sup> is hydrogen, hydroxyl, —NH<sub>2</sub>, —NH(C<sub>1</sub>-C<sub>4</sub>-alkyl), —N(C<sub>1</sub>-C<sub>4</sub>-alkyl)<sub>2</sub>, halogen, C<sub>1</sub>-C<sub>4</sub>-alkyl, or C<sub>1</sub>-C<sub>4</sub>-haloalkyl [C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy or C<sub>1</sub>-C<sub>4</sub>-alkylthio];

X is CR<sup>14</sup>, where R<sup>14</sup> is hydrogen or C<sub>1</sub>-C<sub>5</sub>-alkyl;

R<sup>3</sup> is hydrogen, hydroxyl, —NH<sub>2</sub>, —NH(C<sub>1</sub>-C<sub>4</sub>-alkyl), —N(C<sub>1</sub>-C<sub>4</sub>-alkyl)<sub>2</sub>, halogen, C<sub>1</sub>-C<sub>4</sub>-alkyl, or C<sub>1</sub>-C<sub>4</sub>-haloalkyl [C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy, NH—O—C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-alkylthio or CR<sup>3</sup> is linked to CR<sup>14</sup> as indicated above to give a 5- or 6-membered ring];

R<sup>4</sup> and R<sup>5</sup>, which can be identical or different, are phenyl or naphthyl, which can be substituted by one or more of the

following [radicals] selected from the group consisting of: halogen, nitro, cyano, hydroxyl,  $C_1$ - $C_4$ -alkyl,  $C_1$ - $C_4$ -haloalkyl,  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -haloalkoxy, phenoxy,  $C_1$ - $C_4$ -alkylthio, amino,  $C_1$ - $C_4$ -alkylamino [or] and  $C_1$ - $C_4$ -dialkylamino; or  $R^4$  and  $R^5$  are

phenyl or naphthyl, which are connected together in the ortho position via a direct linkage, a methylene, ethylene or ethenylene group, an oxygen or sulfur atom [or], an  $SO_2$ ,  $NH$  or  $N$ -alkyl group[.] or a  $C_3$ - $C_7$ -cycloalkyl group;

$R^6$  is hydrogen, or  $R^6$  is  $C_1$ - $C_8$ -alkyl,  $C_3$ - $C_6$ -alkenyl,  $C_3$ - $C_6$ -alkynyl or  $C_3$ - $C_8$ -cycloalkyl, where each of these [radicals] can be substituted by one or more [times by]

substituents selected from the group consisting of: halogen, nitro, cyano,  $C_1$ - $C_4$ -alkoxy,  $C_3$ - $C_6$ -alkenyloxy,  $C_3$ - $C_6$ -alkynyoxy,  $C_1$ - $C_4$ -alkylthio,  $C_1$ - $C_4$ -haloalkoxy,

$C_1$ - $C_4$ -alkylcarbonyl,  $C_1$ - $C_4$ -alkoxycarbonyl,  $C_3$ - $C_8$ -alkylcarbonylalkyl,  $C_1$ - $C_4$ -alkylamino, di- $C_1$ - $C_4$ -alkylamino, phenyl [or] and phenoxy which phenyl or phenoxy is substituted by one or more [times by]

substituents selected from the group consisting of: halogen, nitro, cyano,  $C_1$ - $C_4$ -alkyl,  $C_1$ - $C_4$ -haloalkyl,  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -haloalkoxy [or] and  $C_1$ - $C_4$ -alkylthio; or

phenyl or naphthyl, each of which can be substituted by one or more of the following [radicals] selected from

the group consisting of: halogen, nitro, cyano, hydroxyl, amino,  $C_1$ - $C_4$ -alkyl,  $C_1$ - $C_4$ -haloalkyl,  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -haloalkoxy, phenoxy,  $C_1$ - $C_4$ -alkylthio,  $C_1$ - $C_4$ -alkylamino,  $C_1$ - $C_4$ -dialkylamino, dioxomethylene [or] and dioxoethylene; or

a five or six-membered heteroaromatic moiety containing (i) one to three nitrogen atoms, [and/or one sulfur or oxygen atom] (ii) one sulfur atom, (iii) one oxygen atom, (iv) one to three nitrogen atoms and one sulfur atom, or (v) one to three nitrogen atoms and one oxygen atom, which heteroaromatic moiety can carry one or more substituents selected from the group consisting of: one to four halogen atoms [and/or], and one or two of the following [radicals] selected from the group consisting of:  $C_1$ - $C_4$ -alkyl,  $C_1$ - $C_4$ -haloalkyl,

$C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -haloalkoxy,  $C_1$ - $C_4$ -alkylthio, phenyl, phenoxy [or] and phenylcarbonyl, it being possible for the phenyl [radicals] in turn to carry one or more substituents selected from the group consisting of: one to five halogen atoms [and/or], and one to three of the following [radicals] selected from the group consisting of:

$C_1$ - $C_4$ -alkyl,  $C_1$ - $C_4$ -haloalkyl,  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -haloalkoxy and [or]  $C_1$ - $C_4$ -alkylthio[.].

Y is sulfur [or], oxygen or a single bond; and Z is sulfur, oxygen,  $—SO—$  or  $—SO_2—$ .

2. The compound of the formula I as defined in claim 1, wherein X is  $CR^{14}$  and  $R^{14}$  is hydrogen.

3. The compound of the formula I as defined in claim 2, wherein R is  $CO_2H$ .

4. The compound of the formula I as defined in claim 2, wherein  $R^2$  and  $R^3$  each is methoxy.]

5. The compound of the formula I as defined in claim 2, wherein  $R^4$  and  $R^5$  each is phenyl.

6. The compound of the formula I as defined in claim 2, wherein  $R^6$  is  $C_1$ - $C_8$ -alkyl.

7. The compound of the formula I as defined in claim 2, wherein Y is oxygen.

8. The compound of the formula I as defined in claim 2, wherein Z is oxygen or sulfur.

9. The compound of the formula I as defined in claim 8, wherein Z is oxygen.

[10. The compound of the formula I as defined in claim 1, wherein

X is  $CH$ ,

Y is oxygen,

Z is oxygen,

R is  $CO_2H$ ,

$R^2$  is methoxy,

$R^3$  is methoxy,

$R^4$  is phenyl,

$R^5$  is phenyl,

$R^6$  is methyl, ethyl or iso-propyl.]

11. The compound of the formula I as defined in claim 1, wherein R is tetrazole, nitrile or a group



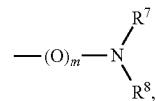
where  $R^1$  has the following meanings:

a) hydrogen;

b) succinylimidoxy;

c) a five-membered heteroaromatic ring linked by a nitrogen atom, selected from the group consisting of: pyrrolyl, pyrazolyl, imidazolyl and triazolyl, which ring can carry one or more substituents selected from the group consisting of: one or two halogen atoms [and/or], and one or two of the following [radicals] selected from the group consisting of:  $C_1$ - $C_4$ -alkyl,  $C_1$ - $C_4$ -haloalkyl,  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -haloalkoxy [or] and  $C_1$ - $C_4$ -alkylthio;

d) a radical



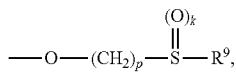
where m is 0 or 1 and  $R^7$  and  $R^8$ , which can be identical or different, have the following meanings:

hydrogen,

$C_1$ - $C_8$ -alkyl,  $C_3$ - $C_6$ -alkenyl,  $C_3$ - $C_6$ -alkynyl,  $C_3$ - $C_8$ -cycloalkyl, where these alkyl, cycloalkyl, alkenyl and alkynyl groups can each carry one or more substituents selected from the group consisting of: one to five halogen atoms [and/or], and one or two of the following groups selected from the group consisting of:  $C_1$ - $C_4$ -alkyl,  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -alkylthio,  $C_1$ - $C_4$ -haloalkoxy,  $C_3$ - $C_6$ -alkenyloxy,  $C_3$ - $C_6$ -alkenylthio,  $C_3$ - $C_6$ -alkynyoxy [or]  $C_3$ - $C_6$ -alkynylthio,  $C_1$ - $C_4$ -alkylcarbonyl,  $C_1$ - $C_4$ -alkoxycarbonyl,  $C_3$ - $C_6$ -alkenylcarbonyl,  $C_3$ - $C_6$ -alkynylcarbonyl,  $C_3$ - $C_6$ -alkenylcarbonyl [or] and  $C_3$ - $C_6$ -alkynylcarbonyl[.]; phenyl, which can be substituted by one or more [times by] substituents selected from the group consisting of: halogen, nitro, cyano,  $C_3$ - $C_6$ -alkenylcarbonyl,  $C_3$ - $C_6$ -alkynylcarbonyl,  $C_1$ - $C_4$ -alkyl,  $C_1$ - $C_4$ -haloalkyl,  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -haloalkoxy [or]  $C_1$ - $C_4$ -alkylthio, and di- $C_1$ - $C_4$ -alkylamino, or

$R^7$  and  $R^8$  together form a  $C_4$ - $C_7$ -alkylene chain which can be substituted by  $C_1$ - $C_4$ -alkyl, and may contain a hetero atom selected from the group consisting of: oxygen, sulfur and nitrogen, or  $R^7$  and  $R^8$  together form a  $CH_2—CH=CH—CH_2$  or  $CH=CH—(CH_2)_3$  chain;

e) a radical



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where k is 0, 1 and 2, p is 1, 2, 3 and 4, and R<sup>9</sup> is C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, C<sub>3</sub>-C<sub>6</sub>-alkenyl, C<sub>3</sub>-C<sub>6</sub>-alkynyl or phenyl, which can be substituted by one or more [times by] *substituents selected from the group consisting of*: halogen, nitro, cyano, C<sub>3</sub>-C<sub>6</sub>-alkenylcarbonyl, C<sub>3</sub>-C<sub>6</sub>-alkynylcarbonyl, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy [or] and C<sub>1</sub>-C<sub>4</sub>-alkylthio;

f) a radical OR<sup>10</sup>, where R<sup>10</sup> is

hydrogen, the cation of an alkali metal or an alkaline earth metal or an environmentally compatible organic ammonium ion;

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C<sub>3</sub>-C<sub>8</sub>-cycloalkyl which may carry one to three C<sub>1</sub>-C<sub>4</sub>-alkyl groups;

C<sub>1</sub>-C<sub>8</sub>-alkyl which may carry *one or more substituents selected from the group consisting of*: one to five halogen atoms [and/or], and one of the following [radicals] *selected from the group consisting of*: C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-alkylthio, cyano, C<sub>1</sub>-C<sub>4</sub>-alkylcarbonyl, C<sub>3</sub>-C<sub>8</sub>-cycloalkyl, C<sub>1</sub>-C<sub>4</sub>-alkoxycarbonyl, phenyl, phenoxy [or] and phenylcarbonyl, where the aromatic [radicals] *substituents* in turn may carry *one or more substituents selected from the group consisting of*: one to five halogen atoms [and/or], and one to three of the following [radicals] *selected from the group consisting of*: nitro, cyano, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy and [or] C<sub>1</sub>-C<sub>4</sub>-alkylthio;

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C<sub>1</sub>-C<sub>8</sub>-alkyl which may carry one to five halogen atoms and which carries one of the following [radicals] *selected from the group consisting of*: a 5-membered heteroaromatic ring containing one to three nitrogen atoms [or], a nitrogen atom and an oxygen [or] and a nitrogen atom and a sulfur atom, which may carry *one or more substituents selected from the group consisting of*: one to four halogen atoms [and/or], and one or two of the following [radicals] *selected from the group consisting of*: nitro, cyano, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, phenyl, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy and [or] C<sub>1</sub>-C<sub>4</sub>-alkylthio;

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C<sub>2</sub>-C<sub>6</sub>-alkyl which carries one of the following [radicals] in position 2: C<sub>1</sub>-C<sub>4</sub>-alkoxyimino, C<sub>3</sub>-C<sub>6</sub>-alkynyl, C<sub>3</sub>-C<sub>6</sub>-alkoxyimino, C<sub>3</sub>-C<sub>6</sub>-haloalkenyl, C<sub>3</sub>-C<sub>6</sub>-alkoxyimino or benzyl, C<sub>3</sub>-C<sub>6</sub>-alkoxyimino;

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C<sub>3</sub>-C<sub>6</sub>-alkenyl or C<sub>3</sub>-C<sub>6</sub>-alkynyl which may carry one to five halogen atoms;

phenyl which may carry *one or more substituents selected from the group consisting of*: one to five halogen atoms [and/or], and one to three of the following [radicals] *selected from the group consisting of*: nitro, cyano, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy and [or] C<sub>1</sub>-C<sub>4</sub>-alkylthio;

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a 5-membered heteroaromatic ring which is bonded via a nitrogen atom and containing one to three nitrogen atoms, which may carry *one or more substituents selected from the group consisting of*: one or two halogen atoms, and [or] one or two of the following [radicals] *selected from the group consisting of*:

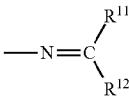
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C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, phenyl, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy and [or] C<sub>1</sub>-C<sub>4</sub>-alkylthio; a radical

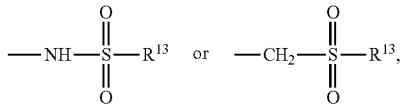


where [R<sup>1</sup>] R<sup>11</sup> and R<sup>12</sup>, which may be identical or different are:

C<sub>1</sub>-C<sub>8</sub>-alkyl, C<sub>3</sub>-C<sub>6</sub>-alkenyl, C<sub>3</sub>-C<sub>6</sub>-alkynyl, C<sub>3</sub>-C<sub>8</sub>-cycloalkyl, it being possible for these [radicals] to carry [a] *one or more substituents selected from the group consisting of*: C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-alkylthio, and [or] phenyl, which may carry *one or more substituents selected from the group consisting of*: one to five halogen atoms [and/or], and one to three of the following [radicals] *selected from the group consisting of*: nitro, cyano, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy and [or] C<sub>1</sub>-C<sub>4</sub>-alkylthio;

phenyl which may carry one or more of the following [radicals] *selected from the group consisting of*: halogen, nitro, cyano, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy [or] and C<sub>1</sub>-C<sub>4</sub>-alkylthio; or R<sup>11</sup> and R<sup>12</sup> together form a C<sub>3</sub>-C<sub>12</sub>-alkylene chain which may carry one to three C<sub>1</sub>-C<sub>4</sub>-alkyl groups and which may contain a hetero atom selected from the group consisting of: a nitrogen, oxygen and sulfur;

g) a radical



where R<sup>13</sup> is

C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>3</sub>-C<sub>6</sub>-alkenyl, C<sub>3</sub>-C<sub>6</sub>-alkynyl, C<sub>3</sub>-C<sub>8</sub>-cycloalkyl, it being possible for these [radicals] to carry *one or more substituents selected from the group consisting of*: a C<sub>1</sub>-C<sub>4</sub>-alkoxy, a C<sub>1</sub>-C<sub>4</sub>-alkylthio [and/or] a phenyl radical, and a phenyl; or

phenyl which may carry one or more of the following [radicals] *selected from the group consisting of*: halogen, nitro, cyano, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy [or] and C<sub>1</sub>-C<sub>4</sub>-alkylthio.

12. The compound of the formula I as defined in claim 1, wherein

R is  $-\text{CO}_2\text{H}$  or a radical which can be hydrolyzed to  $-\text{CO}_2\text{H}$ ;

R<sup>4</sup> is phenyl; and

R<sup>5</sup> is phenyl.

13. The compound of the formula I as defined in claim 12, wherein

X is CH;

Y is oxygen;

Z is oxygen;

R is  $-\text{CO}_2\text{H}$ ;

R<sup>2</sup> is C<sub>1</sub>-C<sub>4</sub>-alkyl;

R<sup>3</sup> is C<sub>1</sub>-C<sub>4</sub>-alkyl; and

R<sup>6</sup> is C<sub>1</sub>-C<sub>8</sub>-alkyl.

14. The compound of the formula I as defined in claim 13, wherein

$R^2$  is methyl; and  
 $R^3$  is methyl.

15. The compound of the formula as defined in claim 1, wherein

$R$  is formyl,  $-CO_2H$  or a radical which can be hydrolyzed to  $-CO_2H$ ;

$R^2$  is  $C_1-C_4$ -alkyl;

$X$  is  $CR^{14}$ , where  $R^{14}$  is hydrogen or  $C_1-C_5$ -alkyl;

$R^3$  is  $C_1-C_4$ -alkyl;

$R^4$  and  $R^5$  which can be identical or different, are phenyl, which can be substituted by one or more of the following selected from the group consisting of: halogen, nitro, hydroxyl,  $C_1-C_4$ -alkyl,  $C_1-C_4$ -alkoxy and  $C_1-C_4$ -alkylthio; or

phenyl which are connected together in the ortho positions by a direct linkage, methylene, ethylene, ethynylene, oxygen, sulfur,  $-SO_2-$ ,  $-NH-$  or  $N$ -alkyl group; or

$C_3-C_7$ -cycloalkyl;

$R^6$  is  $C_1-C_8$ -alkyl,  $C_3-C_6$ -alkenyl or  $C_3-C_8$ -cycloalkyl, where each of these can be substituted by one or more substituents selected from the group consisting of: halogen, hydroxyl, nitro, cyano,  $C_1-C_4$ -alkoxy,  $C_3-C_6$ -alkenyl and  $C_1-C_4$ -alkylthio; or

phenyl or naphthyl, each of which can be substituted by

one or more of the following selected from the group consisting of: halogen, nitro, cyano, hydroxyl, amino,  $C_1-C_4$ -alkyl,  $C_1-C_4$ -haloalkyl,  $C_1-C_4$ -alkoxy,  $C_1-C_4$ -haloalkoxy, phenoxy,  $C_1-C_4$ -alkylthio,  $C_1-C_4$ -alkylamino and  $C_1-C_4$ -dialkylamino; or

a five- or six-membered heteroaromatic moiety containing (i) a nitrogen atom, (ii) a sulfur atom, (iii) an oxygen atom, (iv) a nitrogen atom and a sulfur atom, or (v) a nitrogen atom and an oxygen atom, which heteroaromatic moiety can carry one or more substituents selected from the group consisting of: one to four halogen atoms, and one or two of the following selected from the group consisting of:  $C_1-C_4$ -alkyl,  $C_1-C_4$ -haloalkyl,  $C_1-C_4$ -alkoxy,  $C_1-C_4$ -alkylthio, phenyl, phenoxy and phenylcarbonyl, it being possible for the phenyl in turn to carry one or more substituents selected from the group consisting of: one to five halogen atoms, and one to three of the following selected from the group consisting of:  $C_1-C_4$ -alkyl,  $C_1-C_4$ -haloalkyl,  $C_1-C_4$ -alkoxy and  $C_1-C_4$ -alkylthio;

$Y$  is sulfur, oxygen or a single bond; and

$Z$  is sulfur, oxygen,  $SO$  or  $SO_2$ .

16. The compound of the formula I as defined in claim 15, wherein

$R$  is  $-CO_2H$ ;

$R^2$  is  $C_1-C_4$ -alkyl;

$X$  is  $CR^{14}$ , where  $R^{14}$  is hydrogen;

$R^3$  is  $C_1-C_4$ -alkyl;

$R^4$  and  $R^5$  which can be identical or different, are phenyl, which can be substituted by one or more of the following selected from the group consisting of: halogen, nitro, hydroxyl,  $C_1-C_4$ -alkyl,  $C_1-C_4$ -alkoxy and  $C_1-C_4$ -alkylthio;

$R^6$  is  $C_1-C_8$ -alkyl, which can be substituted by one or more substituents selected from the group consisting of: halogen, hydroxyl, nitro, cyano,  $C_1-C_4$ -alkoxy,  $C_3-C_6$ -alkenyl and  $C_1-C_4$ -alkylthio; or

$Y$  is oxygen; and

$Z$  is oxygen.

17. The compound of the formula I as defined in claim 16, where  $R^4$  and  $R^5$  are each phenyl.

18. The compound of the formula I as defined in claim 1, wherein

$R$  is  $-CO_2H$  or a radical which can be hydrolyzed to  $-CO_2H$ ;

$R^2$  is  $C_1-C_4$ -alkyl;

$X$  is  $CR^{14}$ , where  $R^{14}$  is hydrogen;

$R^3$  is  $C_1-C_4$ -alkyl;

$R^4$  and  $R^5$  which can be identical or different, are phenyl, which can be substituted by one or more of the following selected from the group consisting of: halogen, nitro, hydroxyl,  $C_1-C_4$ -alkyl,  $C_1-C_4$ -alkoxy and  $C_1-C_4$ -alkylthio;

$R^6$  is  $C_1-C_8$ -alkyl, which can be substituted by one or more substituents selected from the group consisting of: halogen, hydroxyl, nitro, cyano,  $C_1-C_4$ -alkoxy,  $C_3-C_6$ -alkenyl and  $C_1-C_4$ -alkylthio;

$Y$  is oxygen; and

$Z$  is oxygen.

19. The compound of the formula I as defined in claim 18, where  $R^4$  and  $R^5$  are each phenyl.

20. The compound of the formula I as defined in claim 1, wherein

$R$  is  $-CO_2H$ ;

$R^2$  is  $C_1-C_4$ -alkyl;

$X$  is  $CR^{14}$ , where  $R^{14}$  is hydrogen or  $C_1-C_5$ -alkyl;

$R^3$  is  $C_1-C_4$ -alkyl;

$R^4$  and  $R^5$  are phenyl which can be substituted by one or more halogen atoms;

$R^6$  is  $C_1-C_8$ -alkyl or  $C_3-C_6$ -cycloalkyl, where each of these can be substituted one or more times by phenyl, or phenyl;

$Y$  is oxygen; and

$Z$  is sulfur or oxygen.

21. The compound of the formula I as defined in claim 1, wherein

$R$  is  $-CO_2H$  or a radical which can be hydrolyzed to  $-CO_2H$ ;

$R^2$  is halogen,  $C_1-C_4$ -alkyl or  $C_1-C_4$ -haloalkyl;

$X$  is  $CR^{14}$ , where  $R^{14}$  is hydrogen or  $C_1-C_5$ -alkyl;

$R^3$  is halogen,  $C_1-C_4$ -alkyl or  $C_1-C_4$ -haloalkyl;

$R^4$  and  $R^5$  are phenyl, which can be substituted by one or more of the following selected from the group consisting of: halogen, nitro,  $C_1-C_4$ -alkyl,  $C_1-C_4$ -haloalkyl,  $C_1-C_4$ -alkoxy,  $C_1-C_4$ -alkylamino and  $C_1-C_4$ -dialkylamino;

$R^6$  is hydrogen, or  $R^6$  is  $C_1-C_8$ -alkyl or  $C_3-C_6$ -cycloalkyl, where each of these can be substituted by one or more substituents selected from the group consisting of: halogen, hydroxyl,  $C_1-C_4$ -alkoxy,  $C_1-C_4$ -alkylthio and phenyl which is substituted by one or more substituents selected from the group consisting of: halogen,  $C_1-C_4$ -alkyl,  $C_1-C_4$ -haloalkyl,  $C_1-C_4$ -alkoxy and  $C_1-C_4$ -alkylthio; or

phenyl which can be substituted by one or more of the following selected from the group consisting of: halogen, nitro, hydroxyl,  $C_1-C_4$ -alkyl,  $C_1-C_4$ -alkoxy,  $C_1-C_4$ -alkylthio and dioxomethylene; or

a five- or six-membered heteroaromatic moiety containing (i) one to three nitrogen atoms, (ii) one sulfur atom, (iii) one oxygen atom, (iv) one to three nitrogen atoms and one sulfur atom, or (v) one to three nitrogen atoms and one oxygen atom which heteroaromatic moiety can carry one to four halogen atoms;

$Y$  is sulfur or oxygen; and

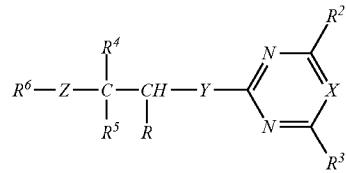
$Z$  is sulfur or oxygen.

22. The compound of claim 21, wherein  
 $R$  is  $-CO_2H$ ,  $-COOCH_3$ ,  $-COON(CH_3)_2$ ,  
 $-COOCH_2C=CH$ ,  $-COOC_2H_5$ ,  $-COON=C(CH_3)_2$ ,  
 $-COONH-phenyl$ ,  $-COOCH=CH_2$  or  
 $-CONH-SO-C_6H_5$ ;  
 $R^2$  is  $-Cl$ ,  $-CH_3$ ,  $-CH_2CH_3$  or  $-CF_3$ ;  
 $X$  is  $CR^{14}$ , where  $R^{14}$  is hydrogen;  
 $R^3$  is  $-Cl$ ,  $-CH_3$ ,  $-CH_2CH_3$  or  $-CF_3$ ;  
 $R^4$  and  $R^5$  are phenyl, which can be substituted by one or  
more groups selected from the group consisting of:  $-F$ ,  
 $-Cl$ ,  $-Br$ ,  $-CH_3$ ,  $-CH_2CH_3$ ,  $-CF_3$ ,  $-OCH_3$ ,  $-NO_2$   
and  $-N(CH_3)_2$ ;  
 $R^6$  is  $-H$ ,  $-CH_3$ ,  $-CH_2CH_3$ ,  $-CH_2CH_2CH_3$ ,  
 $-C(CH_3)_3$ ,  $-CH_2CH(CH_3)_2$ ,  $cyclopropyl$ ,  
 $-CH_2CH_2CH_2CH(CH_3)_2$ ,  $-CH_2-CH_2-S-CH_3$ ,  
 $-CH_2CH_2OH$ , phenyl, trifluoroethyl, *p*-isopropyl-phenyl,  
*p*-methyl-*S*-phenyl, *p*-methyl-*O*-phenyl, *m*-ethyl-phenyl,  
*o*-methyl-phenyl, *o*-Cl-phenyl, *m*-Br-phenyl,  
*p*-F-phenyl, *p*-methyl-phenyl, *m*-NO<sub>2</sub>-phenyl, *o*-HO-phenyl,  
 $3,4$ -dimethoxy-phenyl,  $3,4$ -dioxomethylene-phenyl,  
 $3,4,5$ -trimethoxy-phenyl, benzyl, *o*-Cl-benzyl,  
*m*-Br-benzyl, *p*-F-benzyl, *o*-methyl-benzyl, *m*-ethyl-benzyl or *p*-isopropyl-benzyl;  
 $Y$  is sulfur or oxygen; and  
 $Z$  is sulfur or oxygen.

23. The compound of claim 22, wherein  $R^4$  and  $R^5$  are each phenyl.

24. A compound of the formula I:

(I)



where

$R$  is  $-CO_2H$ ;

$R^2$  is  $C_1-C_4$ -alkyl;

$X$  is  $CR^{14}$ , where  $R^{14}$  is hydrogen;

$R^3$  is  $C_1-C_4$ -alkyl;

$R^4$  and  $R^5$  which can be identical or different, are phenyl, which can be substituted by one or more of the following selected from the group consisting of: halogen, nitro, hydroxyl,  $C_1-C_4$ -alkyl,  $C_1-C_4$ -alkoxy and  $C_1-C_4$ -alkylthio;

$R^6$  is a  $C_1-C_8$ -alkyl, which can be substituted by one or more substituents selected from the group consisting of: halogen, hydroxyl, nitro, cyano,  $C_1-C_4$ -alkoxy,  $C_3-C_6$ -alkenylxy and  $C_1-C_4$ -alkylthio;

$Y$  is oxygen; and

$Z$  is oxygen.

25. The compound of the formula I as defined in claim 24, wherein  $R^4$  and  $R^5$  are each phenyl.

\* \* \* \* \*